

Version 5.1.3  
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using sw model

12, 2002, 01:06:16 ; Search time 286.304 Seconds  
(without alignments)  
1829.698 Million cell updates/sec

US-09-355-254F-10  
1 agctatgacgttccaagg 18  
Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600  
Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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41: em.htgo.other.\*

Pred. No. is the number of results predicted by chance to have a

Score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

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1	18	100.0	18	6	A89789
2	18	100.0	18	6	A90876
3	18	100.0	18	6	AX105148
4	18	100.0	18	6	AX105148 Sequence
5	17	94.4	18	6	AX455555
6	17	94.4	18	6	AX104114
7	17	94.4	18	6	AX353538
8	17	94.4	24	6	AX463126
9	13.2	73.3	37	6	AX463127
10	13.2	73.3	39	6	AR089858
11	13.2	73.3	48	6	AR167631
12	12.8	71.1	19	6	AR151177
13	12.8	71.1	19	6	AX103935
14	12.8	71.1	19	6	AX131449
15	12.8	71.1	41	6	AX353538
16	12.8	71.1	41	6	AR169564
17	12.8	71.1	65	6	AX485786
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19	12.4	68.9	20	6	AR043709
20	12.4	68.9	40	6	AR172301
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27	12.2	67.8	30	6	AX128526
28	12.2	67.8	39	6	A67624
29	12.2	67.8	46	6	AR089762
30	12.2	67.8	49	9	E22537
31	12.2	67.8	98	9	HUMCFDELA
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34	11.8	65.6	25	6	AX355530
35	11.8	65.6	25	6	AX027188
36	11.8	65.6	26	6	AX027200
37	11.8	65.6	35	6	AR105706
38	11.8	65.6	35	6	AX155967
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41	11.8	65.6	40	6	AR182122
42	11.8	65.6	43	3	CCA427110
43	11.8	65.6	51	6	AX162989
44	11.8	65.6	51	6	AX162991
45	11.8	65.6	51	6	AX162993

ALIGNMENTS

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ACCESSION A89789  
VERSION A89789.1  
KEYWORDS GI:6738303  
SOURCE unidentified.  
ORGANISM unclassified.  
REFERENCE 1 (bases 1 to 18)  
AUTHORS Lipford,G.B. and Heeg,K.  
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
JOURNAL OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
Patent: WO 9832462-A 11 30-JUL-1998;

linear PAT 22-JAN-2000

Thu Dec 12 07:53:25 2002

FEATURES  
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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)  
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ACCESSION A90876  
VERSION A90876.1 GI:6739275  
KEYWORDS  
SOURCE  
ORGANISM  
unidentified.  
unclassified.  
REFERENCE 1 (bases 1 to 18)  
AUTHORS Heeg,K.P. and Lipford,G.B.  
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination  
JOURNAL Patent: EP 0855184-A 11 29-JUL-1998;  
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)  
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ACCESSION AX105148  
VERSION AX105148.1 GI:13921298  
KEYWORDS  
SOURCE  
ORGANISM  
synthetic construct.  
synthetic construct.  
artificial sequences.  
REFERENCE 1 (bases 1 to 18)  
AUTHORS Hartmann,G.D., Bratzler,R.L. and Krieg,A.U.  
TITLE Methods related to immunostimulatory nucleic acid-induced interferon  
JOURNAL Patent: WO 0122990-A 46 05-APR-2001;  
Coley Pharmaceutical Group, Inc. (US) ; UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)  
Location/Qualifiers  
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VERSION AX455555.1 GI:21714623  
KEYWORDS  
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artificial sequences.  
REFERENCE 1  
AUTHORS Bauer,S., Lipford,G. and Wagner,H.  
TITLE Process for high throughput screening of cpg-based immuno-agonist/antagonist  
JOURNAL Patent: WO 0222809-A 32 21-MAR-2002;  
Coley Pharmaceutical GmbH (DE)  
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ACCESSION AX104114  
VERSION AX104114.1 GI:13920311  
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artificial sequences.  
REFERENCE 1 (bases 1 to 18)  
AUTHORS Krieg,A.M., Schetter,C. and Vollmer,J.C.  
TITLE Immunostimulatory nucleic acids  
JOURNAL Patent: WO 0122972-A 306 05-APR-2001;  
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical GmbH (DE)  
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TITLE	Screening methods for compounds useful in the regulation of body weight
JOURNAL FEATURES	Patent: US 6287763-A 41 11-SEP-2001; Location/Qualifiers

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DEFINITION Sequence 3 from patent US 6232061.
ACCESSION AR151177
VERSION AR151177.1 GI:15117227
KEYWORDS
SOURCE
ORGANISM Unknown.
REFERENCE
1 (bases 1 to 48)
AUTHORS Marchionni, M. Andrew. and Johnson, C. D.
TITLE Homology cloning.
JOURNAL Patent: US 6232061-A 3 15-MAY-2001;
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Location/Qualifiers
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ACCESSION AX103935
VERSION AX103935.1 GI:13920132
KEYWORDS
SOURCE
ORGANISM synthetic construct.
artificial sequences.
REFERENCE
1 (bases 1 to 19)
AUTHORS Krieg, A. M., Schetter, C. and Vollmer, J. C.
TITLE Immunostimulatory nucleic acids
JOURNAL Patent: WO 0122972-A 127 05-APR-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US) ; Coley Pharmaceutical
GmbH (DE)
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RESULT 15
AR169564 AR169564 41 bp DNA linear PAT 17-DEC-2001
LOCUS
DEFINITION Sequence 62 from patent US 6291173.

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RESULT 13
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LOCUS
DEFINITION Sequence 2667 from Patent W00130362.
ACCESSION AX131449
VERSION AX131449.1 GI:14137754
KEYWORDS
SOURCE
ORGANISM Homo sapiens
human.
REFERENCE
1 (bases 1 to 19)
AUTHORS Robbins, J. M. and Tritz, R.
TITLE Ribozyme therapy for the treatment of proliferative skin and eye
diseases
JOURNAL Patent: WO 0130362-A 2667 03-MAY-2001;
IMMUSOL, INC. (US)
FEATURES
Location/Qualifiers
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ACCESSION AX355385
VERSION AX355385.1 GI:18620053
KEYWORDS
SOURCE
ORGANISM synthetic construct.
artificial sequences.
REFERENCE
1
AUTHORS Weiner, G. and Hartmann, G.
TITLE Methods for enhancing antibody-induced cell lysis and treating
cancer
JOURNAL Patent: WO 0197843-A 413 27-DEC-2001;
UNIVERSITY OF IOWA RESEARCH FOUNDATION (US)
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us-09-355-254f-10.rge

Thu Dec 12 07:53:25 2002

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ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 41)
AUTHORS     Bartel,P.L. and Tavtigian,S.V.
TITLE       WMS2--an MMAC1 interacting protein
JOURNAL     Patent: US 6291173-A 62 18-SEP-2001;
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Search completed: December 12, 2002, 02:55:35
Job time : 293.304 secs

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GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:04:50 ; Search time 102.319 Seconds  
(without alignments)  
440.192 Million cell updates/sec

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Searched: 2185239 seqs, 1125999159 residues

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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

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4	20	100.0	21	19 AAV44856	Probe for AP-1 gen
5	20	100.0	21	19 AAV39812	API sequence A. S
6	20	100.0	21	20 AAX83976	API sequence A. U
7	20	100.0	21	20 AAV82453	API comp oligonucle
8	20	100.0	21	21 AAZ82453	Rabbit AP-1 bindin
9	20	100.0	21	22 AAH26603	AP-1 oligonucleoti

10	20	100.0	21	22	AAH26603	Rat AP-1 synthetic
11	20	100.0	21	22	AAF87560	Consensus binding
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13	18.4	92.0	21	18	AAH85833	API oligonucleotide
14	18.4	92.0	21	20	AAZ25683	Transcription fact
15	18.4	92.0	21	20	AAH60221	Oligonucleotide AP
16	18.4	92.0	21	20	AAH76049	cAMP response elem
17	18.4	92.0	21	20	AAV08337	CRE element coding
18	18.4	92.0	21	21	AAA52331	AP-1 footprinting
19	18.4	92.0	21	22	AAI70580	Transcription fact
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22	18.4	92.0	21	24	ABK97978	Cell-TRAP method a
23	18.4	92.0	21	24	ABK97979	Cell-TRAP method a
24	18.4	92.0	21	24	ABK98231	Nucleic acid bindi
25	18.4	92.0	21	24	ABK98232	Nucleic acid bindi
26	18.4	92.0	21	24	ABL60768	Nuclear factor-kap
27	16.8	84.0	21	19	AAV44855	Probe for AP-1 gen
28	16.8	84.0	21	22	AAH26605	AP-1 mutant oligon
29	16.8	84.0	21	22	AAH13505	Rat AP-1 synthetic
30	15.2	76.0	21	22	AAH13507	Rat AP-1 synthetic
31	15.2	76.0	21	24	ABL60767	Nuclear factor-kap
32	14.4	72.0	19	24	ABL44268	Human chromosome 1
33	14.4	72.0	22	15	AAQ67303	Detection probe fo
34	14.4	72.0	22	21	AAZ56897	AP-1 consensus seq
35	14.4	72.0	25	22	AAF99633	Immunostimulatory
36	14.4	72.0	25	22	AAF99634	Immunostimulatory
37	14.4	72.0	25	24	ABL38910	Immunostimulatory
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42	13.8	69.0	21	22	AAI70282	Phage lambda genom
43	13.8	69.0	25	24	AAH42574	Phospholipase A1 p
44	13.8	69.0	47	21	AAZ68783	Human map-related
45	13.8	69.0	60	22	AAF82009	1.0 kb DNA fragmen

#### ALIGNMENTS

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ID AAV46003 standard; DNA; 20 BP.  
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AC AAV46003;  
XX  
DT 16-OCT-1998 (first entry)  
XX  
DE Immune adjuvant AP-1 #2.  
XX  
KW Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;  
KW modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;  
KW Ig class; autoimmune response; T-cell; B-cell; tumour; ss.  
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OS Class Bacteria.  
XX  
PN EP855184-A1.  
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PD 29-JUL-1998.  
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PF 23-JAN-1997; 97EP-0101019.  
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PR 23-JAN-1997; 97EP-0101019.  
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PA (LIPF/) LIPFORD G B.  
PA (WAGN/) WAGNER H.  
PI Heeg K, Lipford GB, Wagner H;  
XX  
WPI; 1998-389630/34.  
XX

PT Antigenic composition comprises polynucleotide fragment and antigen  
PT - used as vaccine to treat or prevent e.g. cancer or pathogen  
PT infections and to modulate immune response e.g. tolerance break and  
PT regulation of TH1/TH2 cells  
XX  
PS Example 5; Page 8; 28pp; English.  
XX  
CC AAV45993-V46019 are fragments of bacterial polynucleotides which are  
CC used as immune adjuvants for inclusion into vaccines to treat cancer and  
CC for prophylaxis and/or treatment of conditions caused by pathogenic  
CC micro-organisms. The polynucleotide is used for modulation of an immune  
CC response and the modulation is selected from the group break of  
CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig  
CC classes, treatment of autoimmune responses and induction of tolerances.  
CC DNA oligomers are used to enhance the reactivity of immune cells to  
CC viral, bacterial and parasitic antigens, to break tolerance in anergic T  
CC and B cells e.g. against tumour antigens, as adjuvants in vaccination  
CC against tumour-defined antigens and immunostimulatory substances in an  
CC immune response against tumours and to suppress immune reactions of the  
CC innate and acquired immune system. The composition is inexpensive and  
CC stable and does not cause lethal shock, which happens with prior art  
CC bacterial sequences.  
XX  
SQ Sequence 20 BP; 5 A; 5 C; 6 G; 4 T; 0 other;  
Query Match 100.0%; Score 20; DB 19; Length 20;  
Best Local Similarity 100.0%; Pred. No. 0.43;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GCTTGATGACTACGCCGAA 20  
DB 1 GCTTGATGACTACGCCGAA 20  
RESULT 2  
AAZ48028  
ID AAZ48028 standard; DNA; 20 BP.  
XX  
AC AAZ48028;  
XX  
DT 08-MAR-2000 (first entry)  
XX  
DE Immune remodeling inducing CpG oligonucleotide control SEQ ID NO:108.  
XX  
KW Haematopoiesis; regulation; CpG oligonucleotide; phosphorothioate;  
KW immune remodeling; thrombopoiesis; anaemia; immune system; cancer;  
KW immune response; allergic reaction; infectious disease; asthma;  
KW thrombocytopaenia; immunohaemolytic disorder; genetic disorder;  
KW haemoglobinopathy; kidney failure; chronic inflammatory disorder;  
KW rheumatoid arthritis; ss.  
XX  
OS Synthetic.  
XX  
PN WO958118-A2.  
XX  
PD 18-NOV-1999.  
XX  
PF 14-MAY-1999; 99WO-IB01285.  
XX  
PR 14-MAY-1998; 98US-0085516.  
XX  
PR 02-FEB-1999; 99US-0241653.  
XX  
PA (CPGI-) CPG IMMUNOPHARMACEUTICALS GMBH.  
XX  
PA (CPGI-) CPG IMMUNOPHARMACEUTICALS INC.  
XX  
PI Wagner H, Lipford G;  
XX  
DR WPI; 2000-062261/05.  
XX  
PT Use of CpG containing oligonucleotides for, e.g. inducing an  
PT antigen-specific immune response  
XX  
PS Example 1; Page 51; 116pp; English.

XX The present invention describes a method using CpG containing  
CC oligonucleotides (ONs) for regulating immune system remodeling and for  
CC regulating haematopoiesis. The method for inducing an antigen-specific  
CC immune response comprises: (1) administering an ON having a sequence  
CC including at least the formula (I); and (2) exposing the subject to an  
CC antigen at least 3 days after the ON is administered to the subject to  
CC produce an antigen-specific immune response: 5' XICGX2 3' (I), where  
CC the ON = includes at least 8 nucleotides; C and G = unmethylated, and  
CC X1 and X2 = nucleotides. The method can be used for inducing an immune  
CC response against an antigen such as cells, cell extracts, proteins,  
CC polysaccharides, polysaccharide conjugates, lipids, glycolipids,  
CC carbohydrates, viral extracts, viruses, bacteria, fungi, parasites and  
CC allergens. It can be used in a subject at risk of developing cancer or  
CC an allergic reaction. It can also be used for treating an infectious  
CC disease, allergic diseases and asthma, as well as thrombocytopaenia  
CC which is drug-induced, due to an autoimmune disorder such as idiopathic  
CC thrombocytopenic purpura, or resulting from accidental or therapeutic  
CC radiation exposure. It can also be used for treating anaemia such as  
CC drug-induced anaemia, immunohaemolytic disorder, genetic disorders such  
CC as haemoglobinopathy and inherited haemolytic anaemia, inadequate  
CC production of adequate iron stores, chronic disease such as kidney  
CC failure, and chronic inflammatory disorder such as rheumatoid arthritis,  
CC or anaemia resulting from accidental or therapeutic radiation exposure.  
CC AAZ47932 to AAZ48029 represent phosphorothioate CpG oligonucleotides  
XX used in the exemplification of the present invention.  
SQ Sequence 20 BP; 5 A; 5 C; 6 G; 4 T; 0 other;  
Query Match 100.0%; Score 20; DB 21; Length 20;  
Best Local Similarity 100.0%; Pred. No. 0.43;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GCTTGATGACTACGCCGAA 20  
DB 1 GCTTGATGACTACGCCGAA 20  
RESULT 3  
AAL39184  
ID AAL39184 standard; DNA; 20 BP.  
XX  
AC AAL39184;  
XX  
DT 05-SEP-2002 (first entry)  
XX  
DE Murine Toll-like receptor related CpG DNA SEQ ID No 59.  
XX  
KW Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.  
XX  
OS Unidentified.  
XX  
PN WO200222809-A2.  
XX  
PD 21-MAR-2002.  
XX  
PF 17-SEP-2001; 2001WO-US29229.  
XX  
PR 15-SEP-2000; 2000US-233035P.  
XX  
PR 23-JAN-2001; 2001US-263657P.  
XX  
PR 17-MAY-2001; 2001US-291726P.  
XX  
PR 22-JUN-2001; 2001US-300210P.  
XX  
PA (COLE-) COLEY PHARM GMBH.  
XX  
PA Bauer S, Lipford G, Wagner H;  
XX  
DR WPI; 2002-393964/42.  
XX  
PT New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,  
PT useful for identifying species specificity of immunostimulatory nucleic  
PT acid and identifying immunostimulatory nucleic acids  
XX

PS Disclosure: Page 76; 195pp; English.

XX The invention relates to isolated murine Toll-like receptors (TLR)9.

CC TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined

CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or

CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their

CC fragments have an amino acid sequence which is identical to human TLR9,

CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino

CC acid of a murine TLR polypeptide. The isolated nucleic acids of the

CC invention are useful for inhibiting TLR9 signalling activity in a cell.

CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid

CC molecules which interact with a TLR polypeptide or its fragment. The

CC TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The

CC TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9

CC signalling activity of a test compound (that is not a nucleic acid, and

CC is a polypeptide or a part of a combinatorial library of compounds) with

CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for

CC identifying species specificity of an ISNA. The isolated nucleic acids of

CC the invention are useful as probes or primers. This polynucleotide

CC sequence represents DNA relating to the isolated Toll-like receptors of

CC the invention.

XX

SQ Sequence 20 BP; 5 A; 5 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 24; Length 20;

Best Local Similarity 100.0%; Pred. No. 0.43;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20

DB 1 GCTTGATGACTCAGCGGAA 20

|||||

RESULT 4

AAV44856

ID AAV44856 standard; DNA; 21 BP.

XX

AC AAV44856;

XX

DT 21-OCT-1998 (first entry)

XX

DE Probe for AP-1 gene.

XX

KW Entry mediator gene; herpesvirus; HVEM; tumour necrosis factor receptor;

KW gene expression regulator; cellular stress; inflammatory response;

KW lymphocyte activity regulator; autoimmune response; probe; ss.

XX

OS Synthetic.

OS Homo sapiens.

XX

PN WO9825967-A1.

XX

PD 18-JUN-1998.

XX

PF 05-DEC-1997; 97WO-US22278.

XX

PR 12-DEC-1996; 96US-0032705.

XX

PA (GETH ) GENENTECH INC.

XX

PI Ashkenazi AJ, Marsters SA;

XX

DR WPI; 1998-348457/30.

XX

PT Herpesvirus entry mediator polypeptide, HVEM - useful, e.g. in

PT assays for HVEM and to produce antibodies and transgenic animals,

PT e.g. for drug screening

XX

PS Example 3; Page 32; 46pp; English.

XX

CC This sequence is a probe for AP-1, used to determine if the

CC herpesvirus entry mediator (HVEM) protein of the invention activates

CC NF-kappaB. The HVEM protein is useful in quantitative diagnostic assays

for HVEM, in affinity purification of HVEM from recombinant cells/natural

sources and in competitive-type receptor binding assays. It can be used

to generate antibodies, also useful in diagnostic assays for HVEM and

affinity purification of HVEM. HVEM is believed to be a member of the

tumour necrosis factor receptor family. Transient transfection of HVEM

into human 293 cells caused marked activation of certain transcription

factors, suggesting that HVEM is involved in regulating gene expression

in response to infectious stimuli and cellular stress. The predominant

expression of HVEM mRNA in lymphocyte-rich tissues (e.g. spleen and

peripheral blood) suggests it may be a receptor in regulating lymphocyte

activity. Antibodies produced may be useful therapeutically, e.g.

antagonistic antibodies may be used to block excessive

inflammatory/autoimmune response resulting from e.g. AP-1 induction,

whilst agonistic antibodies may enhance HVEM regulation of such

induction. The DNA may be used diagnostically, e.g. to determine if DNA

and/or RNA encoding HVEM is present in cells, and to prepare HVEM

polypeptide recombinantly. It is also useful to produce non-human

transgenic animals (e.g. mice or rats), especially knockout animals

containing cells with an altered gene encoding HVEM polypeptide. Such

animals are useful in the development and screening of therapeutically

useful reagents.

XX

SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 19; Length 21;

Best Local Similarity 100.0%; Pred. No. 0.43;

Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20

DB 2 GCTTGATGACTCAGCGGAA 21

|||||

RESULT 5

AAV39812

ID AAV39812 standard; DNA; 21 BP.

XX

AC AAV39812;

XX

DT 24-SEP-1998 (first entry)

XX

DE AP1 sequence A.

XX

KW Mouse type 10 collagen promoter; AP1 sequence A; mutation; MEF-2;

KW identification; morphogen; Op-1; osteogenic protein 1; human c-fos; ss.

XX

OS Synthetic.

OS Homo sapiens.

XX

PN WO9826069-A1.

XX

PD 18-JUN-1998.

XX

PF 11-DEC-1997; 97WO-US23211.

XX

PR 12-DEC-1996; 96US-0764522.

XX

PA (CREA-) CREATIVE BIOMOLECULES INC.

XX

PI Harada S, Rodan GA, Sampath KT;

XX

DR WPI; 1998-348526/30.

XX

PT Identifying morphogen analogues - by using DNA defining a

PT morphogen-responsive transcription activating element from a mouse

PT type X collagen promoter

XX

PS Claim 10; Page 61; 88pp; English.

XX

CC A method has been developed for identifying a compound that induces a

CC morphogen-mediated biological effect. The method comprises: (a) providing

CC a test cell comprising DNA defining a morphogen-responsive transcription

CC activating element (MRTAE), and, in operative association, a reporter

CC gene encoding a detectable gene product, the DNA, when present in a  
 CC morphogen-responsive cell contacted with morphogen, serving to induce  
 CC transcription of the reporter gene; (b) exposing the test cell to a  
 CC candidate compound, and (c) detecting expression of the detectable gene  
 CC product, the expression indicating the ability of the candidate compound  
 CC to induce the morphogen-mediated biological effect. The present invention  
 CC also describes: (1) a method for assessing whether a sample comprises a  
 CC substance competent to bind to DNA; (2) a method for identifying a  
 CC candidate compound that induces a morphogen-mediated biological effect;  
 CC (3) a method for monitoring cell differentiation or tissue morphogenesis;  
 CC and (4) a method for identifying a tissue responsive to a morphogen or  
 CC analogue. The methods can be used for obtaining morphogen analogues. In  
 CC particular they can be used for obtaining analogues of osteogenic protein  
 CC 1 (OP-1) for the treatment of a metabolic bone disease, e.g. a disease  
 CC characterised by osteopenia. Analogues can also be obtained for treating  
 CC mammals afflicted with ischemic, ulcerative or inflammatory tissue  
 CC damage, or with injury or deterioration of a morphogen-responsive tissue  
 CC such as bone, liver, kidney, nerve, gastrointestinal tract lining, tooth  
 CC dentin or periodontal tissue. The present sequence represents API  
 CC sequence A from the present invention.

XX Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 19; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 0.43;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 |||||  
 Db 2 GCTTGATGACTCAGCGGAA 21

# RESULT 6

AA83976  
 ID AA83976 standard; DNA; 21 BP.

XX AC AA83976;

XX 08-SEP-1999 (first entry)

DE API sequence A.

XX Mouse; type 10 collagen promoter; API sequence A; osteogenic protein;  
 KW OP-1; morphogen; bone morphogenic protein; BMP; soft tissue disorder;  
 KW apoptosis; morphogen-activated regulatory pathway; tumour;  
 KW cellular immune rejection; viral disease; ss.

XX Unidentified.

XX WO9931136-A2.

XX 24-JUN-1999.

XX 16-DEC-1998; 98WO-US26788.

XX 01-DEC-1998; 98US-0110498.

XX 17-DEC-1997; 97US-0069931.

XX (CREA-) CREATIVE BIOMOLECULES INC.

XX Cohen CM, Kawabata M, Miyazono K, Oeda E, Sampath KT;

XX WPI; 1999-418756/35.

XX Maintaining or restoring tissue-appropriate phenotype

XX Disclosure; Page 45; 50pp; English.

XX A method has been developed for maintaining or restoring tissue-  
 CC appropriate phenotype by expression of a phenotype-specific protein or  
 CC by inhibiting an intracellular pathway that induces expression of a  
 CC gene that is an inhibitor of normal phenotype. The method is for  
 CC restoring cellular phenotype in a cell effected by disease, damage or

CC age. The method comprises activating an intracellular pathway that  
 CC induces expression of a phenotype-specific gene. Another method is also  
 CC described for restoring cellular phenotype in a cell effected by  
 CC disease, damage or age, comprising inhibiting an intracellular pathway  
 CC that induces expression of a gene (especially TGF-beta) that is an  
 CC inhibitor of normal phenotype. The methods can be used to treat soft  
 CC tissue disorders by affecting apoptosis by modulating a morphogen-  
 CC activated regulatory pathway e.g. in tumours, cellular immune rejection  
 CC and viral diseases. The present sequence is used in the exemplification  
 CC of the present invention.

XX Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 0.43;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 |||||  
 Db 2 GCTTGATGACTCAGCGGAA 21

# RESULT 7

AAV82453  
 ID AAV82453 standard; DNA; 21 BP.

XX AC AAV82453;

XX 12-APR-1999 (first entry)

DE API comp oligonucleotide used in competition analysis.

KW Vascular endothelial growth factor; VEGF; human; hypoxia;  
 KW vascular disease; tumour; cancer; angiogenesis; wound healing;  
 KW therapy; diagnosis; ds.

XX Synthetic.

XX Homo sapiens.

XX WO9856936-A1.

XX 17-DEC-1998.

XX 10-JUN-1998; 98WO-EP03517.

XX 10-JUN-1997; 97EP-0109418.

XX (PLAC ) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.

XX Damert A, Plate K, Risau W;

XX WPI; 1999-080911/07.

XX New recombinant DNA - contains sequence that regulates  
 PT hypoxia-induced expression, used for, e.g. treatment and diagnosis  
 PT of vascular disease

XX Example 6; Page 41; 80pp; English.

XX Oligonucleotides hVEGF, hVEGF 5' DEL, API1 and API2 (see  
 CC AAV82449-52), and competitor oligonucleotides API comp, ATF comp  
 CC and VL30 (see AAV82453-55) were used in electrophoretic mobility  
 CC shift assays to determine which transcription factor(s) bind to  
 CC the cis-acting element that is involved in the potentiation of  
 CC hypoxia inducible factor 1 (HIF-1) mediated hypoxic induction  
 CC of vascular endothelial growth factor (VEGF) gene regulatory  
 CC sequences. Experiments were performed using normoxic or hypoxic  
 CC C6 cell nuclear extracts. An API consensus binding site was shown  
 CC to compete for DNA-protein-complex formation at potentiating  
 CC sequences. The invention relates to recombinant DNA molecules  
 CC comprising regulatory sequences of the VEGF gene, especially the  
 CC 3' untranslated region (see AAV82439) and promoter (see AAV82440),  
 CC being capable of modulating hypoxia inducible expression of a

CC heterologous DNA in vivo. Such recombinant DNA molecules, vectors,  
CC host cells and transgenic animals can be used to identify and  
CC develop compounds and methods for diagnosing, treating, preventing  
CC and/or delaying a vascular or tumour disease.  
XX  
SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;  
  
Query Match 100.0%; Score 20; DB 20; Length 21;  
Best Local Similarity 100.0%; Pred. No. 0.43;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 1 GCTTGATGACTCAGCCGGAA 20  
DB 2 GCTTGATGACTCAGCCGGAA 21  
|||||  
  
RESULT 8  
AAZ89651  
ID AAZ89651 standard; DNA; 21 BP.  
XX  
AC AAZ89651;  
CC  
DT 28-JUN-2000 (first entry)  
XX  
DE Rabbit AP-1 binding site, consensus DNA sequence.  
XX  
KW Transcription factor; AP-1; C/EBP; CCAAT/enhancer binding protein;  
KW cardiant; gene therapy; coronary angioplasty; stent implantation;  
KW ET-1 gene; smooth muscle cell; aorta carotis; treatment; rabbit;  
KW coronary heart disease; ds.  
XX  
OS Oryctolagus cuniculus.  
XX  
PN DE29916160-U1.  
XX  
PD 09-MAR-2000.  
XX  
PF 31-AUG-1999; 99WO-US20047.  
XX  
PR 25-MAY-1999; 99DE-1023892.  
XX  
PA (CARD-) CARDIOGENE GENTHERAPEUTISCHE SYSTEME AG.  
XX  
DR WPI; 2000-247751/22.  
XX  
CC Double stranded oligonucleotides targeting genes encoding AP-1, C/EBP  
PT and related transcription factors useful for treatment of coronary  
PT heart disease -  
XX  
PS Example 6; Page 37; 53pp; German.  
XX  
CC This invention describes novel double stranded nucleic acids, which bind  
CC specifically to transcription factors AP-1, C/EBP (CCAAT/Enhancer  
CC Binding Protein) or related transcription factors. The products of the  
CC invention have cardiant activity and are used in gene therapy. Patients  
CC who have had coronary angioplasty or stent implantation may be at risk  
CC from an increased expression of certain genes due to the physical weight  
CC of the devices used inducing a number of genes involved in the cell  
CC cycle. The AP-1 and C/EBP specific double stranded nucleic acids can be  
CC used to block activation of the ET-1 gene in smooth muscle cells of the  
CC aorta carotis. The nucleic acids are therefore useful in treatment of  
CC coronary heart diseases. This sequence represents a consensus AP-1  
CC binding site isolated from rabbit (Oryctolagus cuniculus) which is used  
CC to illustrate the method of the invention.  
XX  
SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;  
  
Query Match 100.0%; Score 20; DB 21; Length 21;  
Best Local Similarity 100.0%; Pred. No. 0.43;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 1 GCTTGATGACTCAGCCGGAA 20  
|||||

Db 2 GCTTGATGACTCAGCCGGAA 21  
  
RESULT 9  
AAH26603  
ID AAH26603 standard; DNA; 21 BP.  
XX  
AC AAH26603;  
XX  
DT 12-NOV-2001 (first entry)  
XX  
DE AP-1 oligonucleotide.  
XX  
KW Melanoma differentiation associated gene-7; Mda-7; human;  
KW promoter; neuroblastoma; astrocytoma; glioblastoma multiforme;  
KW cervical cancer; breast cancer; colon cancer; prostate cancer;  
KW osteosarcoma; chondrosarcoma; tumour; therapy; PCR primer;  
KW electrophoretic mobility shift assay; EMSA; AP-1;  
KW transcription factor; ss.  
XX  
OS Homo sapiens.  
XX  
PN WO200164921-A1.  
XX  
PD 07-SEP-2001.  
XX  
PF 28-FEB-2001; 2001WO-US06782.  
XX  
PR 29-FEB-2000; 2000US-0515369.  
XX  
PA (UYCO ) UNIV COLUMBIA NEW YORK.  
XX  
PI Fisher PB, Madireddi MT;  
XX  
DR WPI; 2001-565508/53.  
XX  
PT Melanoma differentiation associated gene-7 promoter capable of  
PT treating cancer comprises directing transcription of heterologous  
PT coding sequence encoding tumour suppressor polypeptide positioned  
PT downstream, useful for treating cancer -  
XX  
PS Example 2; Page 70; 132pp; English.  
XX  
CC The present sequence is that of an AP-1 transcription factor  
CC oligonucleotide. This was used in electrophoretic mobility  
CC shift assays (EMSA) to examine transcription factor binding to  
CC the human melanoma differentiation associated gene-7 (mda-7)  
CC promoter (see AAH26595). Results demonstrated that cJUN/AP-1 and  
CC C/EBP-beta transcription factors bind to defined sites within the  
CC mda-7 promoter during the process of terminal differentiation in  
CC human melanoma cells. The invention provides recombinant  
CC expression constructs in which the mda-7 promoter is operably  
CC linked to a coding sequence encoding a tumour suppressor protein.  
CC A pharmaceutical composition including the recombinant expression  
CC construct is used in a claimed method of treating melanoma,  
CC neuroblastoma, astrocytoma, glioblastoma multiforme, cervical  
CC cancer, breast cancer, colon cancer, prostate cancer, osteosarcoma,  
CC chondrosarcoma or a cancer of the central nervous system.  
XX  
SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;  
  
Query Match 100.0%; Score 20; DB 22; Length 21;  
Best Local Similarity 100.0%; Pred. No. 0.43;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 1 GCTTGATGACTCAGCCGGAA 20  
|||||  
Db 2 GCTTGATGACTCAGCCGGAA 21  
  
RESULT 10  
AAD13504  
ID AAD13504 standard; DNA; 21 BP.

XX AAD13504;  
 XX 06-NOV-2001 (first entry)  
 XX Rat AP-1 synthetic duplex probe #1.  
 XX  
 KW Antioxidative enzyme; catalase; CAT; superoxide dismutase; SOD; therapy;  
 KW reactive oxygen species; ROS; free radical; dietary supplement; stroke;  
 KW AP-1 transcription factor; renal reperfusion damage; cerebral ischaemia;  
 KW myocardial infarction; heart attack; pain; atherosclerosis; neuroleptic;  
 KW trauma; premature aging; neurodegenerative disease; tardive dyskinesia;  
 KW Parkinson's disease; amyotrophic lateral sclerosis; Alzheimer's disease;  
 KW arthritis; inflammatory disease; diabetes; ulcerative colitis; cataract;  
 KW senility; Down's syndrome; macular degeneration; septic shock; epilepsy;  
 KW polytraumatic shock; schizophrenia; antilucer; clozapine; tranquilliser;  
 KW cardiac; cerebroprotective; vulnerary; nootropic; Huntington's disease;  
 KW anticonvulsant; neuroprotective; antiarthritic; antiinflammatory; burn;  
 KW cytostatic; leukaemia; ophthalmological; antibacterial; probe; ds.  
 XX  
 OS Rattus sp.  
 XX  
 PN WO200136454-A1.  
 XX  
 XX 25-MAY-2001.  
 XX  
 PF 17-NOV-2000; 2000WO-US31764.  
 XX  
 PR 18-NOV-1999; 99US-0166381.  
 XX  
 XX (CERE-) CEREMEDIX INC.  
 XX  
 XX Shashoua VE;  
 XX  
 XX WPI: 2001-496512/54.  
 DR  
 XX  
 PT Novel peptide compound that up regulates expression of a gene encoding  
 PT antioxidative enzymes, used to treat or prevent conditions caused by  
 PT undesirable elevation of reactive oxygen species and other free  
 PT radicals -  
 XX  
 PS Example 3; Page 50; 102pp; English.  
 XX  
 CC The invention relates to peptide compounds and methods for upregulating  
 CC expression of a gene encoding an antioxidative enzyme, such as catalase  
 CC (CAT) or superoxide dismutase (SOD), to counteract harmful oxidative  
 CC effects of reactive oxygen species (ROS) and other free radicals. The  
 CC peptides are used as components of pharmaceuticals and dietary  
 CC supplements. The peptides are used to treat or to prevent diseases and  
 CC conditions characterised by undesirable elevation of ROS and other free  
 CC radicals, to upregulate AP-1 transcription factor gene expression and to  
 CC treat pain. The disease or conditions include renal reperfusion damage,  
 CC cerebral ischaemia (stroke), myocardial infarction (heart attack), head  
 CC trauma, atherosclerosis, brain trauma, oxygen toxicity in premature  
 CC infants, premature aging, spinal cord trauma, neurodegenerative diseases,  
 CC Huntington's disease, Parkinson's disease, amyotrophic lateral sclerosis,  
 CC Alzheimer's disease, arthritis and other inflammatory diseases, diabetes,  
 CC ulcerative colitis, human leukaemia and other cancers characterised by  
 CC elevation of ROS or free radicals, age-related elevation of ROS or free  
 CC radicals, senility, Down's syndrome, macular degeneration, cataracts,  
 CC septic shock, polytraumatic shock, schizophrenia, burn injuries,  
 CC epilepsy, radiation and/or drug-induced elevation of ROS and free  
 CC radicals, where the drug is a neuroleptic or a drug such as clozapine  
 CC defined in the specification and Tardive dyskinesia. The present  
 CC sequence is a duplex probe used to assay rat AP-1 transcription  
 CC factor activation.  
 XX  
 SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;  
 Query Match 100.0%; Score 20; DB 22; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 0.43;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 |||||  
 Db 2 GCTTGATGACTCAGCGGAA 21  
 RESULT 11  
 AAF87560  
 ID AAF87560 standard; DNA; 21 BP.  
 XX  
 AC AAF87560;  
 XX  
 DT 04-JUL-2001 (first entry)  
 XX  
 DE Consensus binding site for AP-1.  
 KW Transcription modulation; AP-1; C/EBP; vascular; coronary; myocardial;  
 KW CCAAT/enhancer binding protein; ds.  
 XX  
 OS Unidentified.  
 XX  
 PN CA2300328-A1.  
 XX  
 PD 14-MAR-2001.  
 XX  
 PF 13-MAR-2000; 2000CA-2300328.  
 XX  
 PR 14-SEP-1999; 99JP-0261035.  
 PR 14-SEP-1999; 99DE-2016160.  
 XX  
 XX (CARD-) CARDIOGENE GENTHERAP SYSTEME AG.  
 XX  
 XX Hecker M, Lauth M, Wagner AH;  
 XX  
 XX WPI: 2001-316865/34.  
 XX  
 PT Modulating gene transcription in an endothelial, vascular or cardiac  
 PT cell, useful for treating vascular proliferative diseases, comprises  
 PT contacting the cell with a nucleic acid capable of sequence-specific  
 PT binding to AP-1 or C/EBP -  
 XX  
 XX Claim 27; Page 36; 57pp; English.  
 XX  
 CC The present invention relates to modulating the transcription of  
 CC at least one gene in an endothelial, vascular or cardiac cell  
 CC by contacting the cell with a composition of one or more double  
 CC stranded nucleic acids capable of sequence-specific binding to  
 CC the transcription factor AP-1 or C/EBP (CCAAT/enhancer binding  
 CC protein). The method is also useful for treating vascular  
 CC diseases, specifically vascular proliferative disorders,  
 CC coronary heart diseases and myocardial infarction. The present  
 CC sequence is the consensus binding site for AP-1.  
 XX  
 SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;  
 Query Match 100.0%; Score 20; DB 22; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 0.43;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 |||||  
 Db 2 GCTTGATGACTCAGCGGAA 21  
 RESULT 12  
 ABL56586  
 ID ABL56586 standard; DNA; 21 BP.  
 XX  
 AC ABL56586;  
 XX  
 DT 30-JUL-2002 (first entry)  
 XX  
 XX Oligonucleotide specific for nuclear transcription factor AP-1.  
 XX



KW Nuclear transcription factor AP-1; R-flurbiprofen; S-flurbiprofen;  
 KW R-arylpropionic acid; joint destruction; antirheumatic;  
 KW rheumatic disease; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO200224190-A1.  
 XX  
 PD 28-MAR-2002.  
 XX  
 PF 24-SEP-2001; 2001WO-EP11004.  
 XX  
 PF 25-SEP-2000; 2000DE-1047319.  
 PR  
 XX  
 XX (PAZA-) PAZ ARZNEIMITTEL ENTWICKLUNGS GMBH.  
 XX  
 XX Geisslinger G, Groesch S;  
 XX  
 XX WPI; 2002-401951/43.  
 DR  
 XX Use of R-arylpropionic acids in the production of medicaments, useful  
 PT for the treatment of illnesses, e.g. rheumatic diseases, influenced by  
 PT activation of nuclear transcription factor -  
 XX  
 XX Example; Page 10; 20pp; German.  
 PS  
 XX  
 CC The present sequence represents an oligonucleotide specific for nuclear  
 CC transcription factor AP-1. The oligonucleotide was used to demonstrate  
 CC that while R-flurbiprofen inhibited AP-1 DNA binding in a  
 CC dosage-dependent manner (binding was completely suppressed at 1000  
 CC microgram), S-flurbiprofen reduced AP-1 DNA binding at only 1000  
 CC microgram). The specification describes the use of R-arylpropionic acids  
 CC or their salts or derivatives in the production of medicaments which  
 CC inhibit the activation of nuclear transcription factor AP-1 and are  
 CC useful for the treatment of illnesses which are influenced by this  
 CC factor. The R-arylpropionic acids arrest joint destruction but do not  
 CC have the severe side effects associated with standard therapy using  
 CC long-term antirheumatics. The medicaments are used in the treatment of  
 CC rheumatic diseases, preferably in combination with basic therapeutics.  
 XX  
 SQ Sequence 21 BP; 5 A; 6 C; 6 G; 4 T; 0 other;  
 XX  
 Query Match 100.0%; Score 20; DB 24; Length 21;  
 Best Local Similarity 100.0%; Pred. No. 0.43;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GCTTGATGACTCAGCCGGAA 20  
 Db | | | | | | | | | | | | | | | | | | | |  
 2 GCTTGATGACTCAGCCGGAA 21  
 RESULT 13  
 AAT85833  
 ID AAT85833 standard; DNA; 21 BP.  
 AC  
 XX  
 AC AAT85833;  
 XX  
 DT 21-NOV-1997 (first entry)  
 XX  
 DE AP1 oligonucleotide used in gel shift assay.  
 XX  
 KW Activating transcription factor 1; ATF1; CREB; recognition sequence;  
 KW cyclic AMP responsive element binding protein; inhibition; binding;  
 KW proliferation; virus; cancer; HTLV1; leukaemia; antibody; ss.  
 XX  
 OS Synthetic.  
 XX  
 XX US5641486-A.  
 PN  
 XX  
 PD 24-JUN-1997.  
 XX  
 PF 18-MAR-1994; 94US-0210880.  
 XX  
 XX

PR 18-MAR-1994; 94US-0210880.  
 XX  
 PA (UYNE-) UNIV NEBRASKA.  
 XX  
 PI Hinrichs SH, Orten DJ;  
 XX  
 XX WPI; 1997-340900/31.  
 DR  
 XX  
 PT Inhibiting replication of cancer cells or viruses - with inhibitor  
 PT that binds to peptide sequence of activating transcription factor 1  
 XX  
 PS Example 2; Column 6; 17pp; English.  
 XX  
 CC This oligonucleotide sequence corresponds to the recognition sequence AP1  
 CC to which members of the activating transcription factor 1 (ATF1)-cyclic  
 CC AMP responsive element binding protein (CREB) family of proteins bind.  
 CC The sequence was used in a gel shift mobility assay to identify agents  
 CC which inhibit the binding of ATF1 to its recognition sequence. The  
 CC agents are preferably antibodies, small molecules or polypeptides,  
 CC especially the complementarity determining region of monoclonal antibody  
 CC MAb4. The agents cause inhibition of transcription by dissociating ATF1  
 CC from its target gene and thus will prevent proliferation of e.g. a virus  
 CC or cancer cell, such as HTLV1-mediated leukaemic cell proliferation.  
 XX  
 SQ Sequence 21 BP; 5 A; 5 C; 7 G; 4 T; 0 other;  
 XX  
 Query Match 92.0%; Score 18.4; DB 18; Length 21;  
 Best Local Similarity 95.0%; Pred. No. 3.1;  
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;  
 QY 1 GCTTGATGACTCAGCCGGAA 20  
 Db | | | | | | | | | | | | | | | | | | | |  
 2 GCTTGATGACTCAGCCGGAA 21  
 RESULT 14  
 AAZ25683  
 ID AAZ25683 standard; DNA; 21 BP.  
 XX  
 AC AAZ25683;  
 XX  
 DT 06-JAN-2000 (first entry)  
 XX  
 DE Transcription factor AP-1 (c-jun) oligonucleotide.  
 XX  
 KW Neuroprotective; calcium binding; stroke; neurodegenerative disease;  
 KW blood-brain barrier; cerebral ischaemia; Alzheimer's disease;  
 KW memory deficit; aging; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN WO9940112-A1.  
 PN  
 XX  
 PD 12-AUG-1999.  
 XX  
 PF 28-JAN-1999; 99WO-US01786.  
 XX  
 XX 10-FEB-1998; 98US-0021247.  
 PR  
 XX  
 PA (NEUR-) NEUROMEDICA INC.  
 XX  
 XX Shashoua VE;  
 PI  
 XX WPI; 1999-610582/52.  
 DR  
 XX  
 PT Neuroprotective peptides, which bind calcium, are useful for treating  
 PT stroke and other neurodegenerative diseases -  
 XX  
 PS Example 4; Page 31; 65pp; English.  
 XX  
 CC The present invention describes a composition comprising an isolated  
 CC peptide, which comprises the amino acid sequence (I) or (Ia):  
 CC X1-XX3-XX5-XX7-XX9-XXX12 (I); X5-X6-X7-X8-X9-X10-X11-X12 (Ia); where

CC X1 = Asp, Gln, Gly or Tyr; X = any amino acid; X3 = Asp, Asn, Thr or Glu;  
 CC X5 = Asp, Ser, Gly, Asn or Leu; X7 = Ala, Asp, Phe, Lys, Thr, Tyr, Arg,  
 CC Val, Cys or Ser; X9 = Asp, Glu, Gly, Ser, Thr, Met or Asn; and X12 =  
 CC Glu, Gln, Ala, Leu or Asn. (I) and (Ia) are neuroprotective calcium  
 CC binding peptides. (I) is used to treat a condition characterized by  
 CC cerebral ischaemia. (I) reduces the neurotoxic effect of cerebral  
 CC ischaemia. (I) is used to increase neuronal cell AP-1 or NF-IL6  
 CC transcription factor activity. The peptides are also useful for binding  
 CC calcium. The peptide can be conjugated with a compound which facilitates  
 CC transport across the blood brain barrier into the brain or it can be  
 CC administered with a compound that increases transport across the blood  
 CC brain barrier. Molecules that protect neurons against the ischaemic  
 CC effects of stroke will also be useful for treating Alzheimer's disease,  
 CC as well as the memory deficits that are characteristic of the aging  
 CC process. The present sequence represents an oligonucleotide used in  
 CC the exemplification of the present invention.  
 XX  
 SQ Sequence 21 BP; 5 A; 5 C; 7 G; 4 T; 0 other;

Query Match 92.0%; Score 18.4; DB 20; Length 21;  
 Best Local Similarity 95.0%; Pred. No. 3.1;  
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCCGGAA 20  
 ||||| ||||| ||||| |||||  
 Db 2 GCTTGATGAGTCAGCCGGAA 21

RESULT 15  
 AAX60221  
 ID AAX60221 standard; DNA: 21 BP.

XX AC AAX60221;

DT 11-AUG-1999 (first entry)

DE Oligonucleotide AP-1.

XX keratin K1 based expression vector; epidermal cell expression;  
 KW mammalian K1 Keratin gene; primer; ss.

XX Synthetic.

XX US5914265-A.

XX 22-JUN-1999.

PF 01-NOV-1993; 93US-0147777.

XX 01-NOV-1993; 93US-0147777.

PR 30-APR-1992; 92US-0876289.

PR 29-OCT-1993; 93US-0145387.

XX (BAYU ) BAYLOR COLLEGE MEDICINE.

PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.

XX Greenhalgh DA, Roop DR, Rothnagel JA, Yuspa SH;

XX WPI; 1999-370501/31.

XX Keratin K1 expression vectors

XX Example 16; Column 35; 50pp; English.

CC The specification describes a keratin K1 based expression vector, which  
 CC does not comprise a K1 keratin encoding gene sequence, for expression  
 CC in an epidermal cell. The vector comprises a 5' flanking region from a  
 CC mammalian gene including the necessary sequences for expression, a  
 CC 3' flanking region from a mammalian K1 keratin gene which regulates  
 CC expression in an epidermal cell, and a linker connecting the 5' flanking  
 CC region to the nucleic acid sequence, where the linker has a position  
 CC for inserting the nucleic acid sequence but lacks the coding sequence  
 CC of a gene with which it is naturally associated. The genetic material

CC comprising the vector may encode a hormone, a receptor, a growth factor,  
 CC an enzyme, a drug, a tumour suppressor, an apolipoprotein, a clotting  
 CC factor, an antigen, an oncogene or a transforming gene. The present  
 CC oligonucleotide was used in the course of the invention.

XX SQ Sequence 21 BP; 5 A; 5 C; 7 G; 4 T; 0 other;

Query Match 92.0%; Score 18.4; DB 20; Length 21;  
 Best Local Similarity 95.0%; Pred. No. 3.1;  
 Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCCGGAA 20  
 ||||| ||||| ||||| |||||  
 Db 2 GCTTGATGAGTCAGCCGGAA 21

Search completed: December 12, 2002, 01:36:26  
 Job time : 105.319 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2002 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds  
(without alignments)  
1829.698 Million cell updates/sec

Title: US-09-355-254F-12

Perfect score: 20

Sequence: 1 tcgatggggcgagcgcgagc 20

Scoring table: IDENTITY\_NUC

Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

GenEmbl.\*

1: gb.ba.\*  
2: gb.htg.\*  
3: gb.in.\*  
4: gb.om.\*  
5: gb.ov.\*  
6: gb.pat.\*  
7: gb.ph.\*  
8: gb.pl.\*  
9: gb.pr.\*  
10: gb.roi.\*  
11: gb.sts.\*  
12: gb.sy.\*  
13: gb.un.\*  
14: gb.vi.\*  
15: em.ba.\*  
16: em.fun.\*  
17: em.in.\*  
18: em.in.\*  
19: em.mu.\*  
20: em.mu.\*  
21: em.or.\*  
22: em.ov.\*  
23: em.pat.\*  
24: em.ph.\*  
25: em.pl.\*  
26: em.ro.\*  
27: em.sts.\*  
28: em.un.\*  
29: em.vi.\*  
30: em.htg\_hum.\*  
31: em.htg\_inv.\*  
32: em.htg\_other.\*  
33: em.htg\_mus.\*  
34: em.htg\_pln.\*  
35: em.htg\_rod.\*  
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41: em.htgo\_other.\*

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	20	100.0	20	6	A89791	A89791 Sequence 13
2	20	100.0	20	6	A90878	A90878 Sequence 13
3	20	100.0	20	6	AX023404	AX023404 Sequence
4	20	100.0	20	6	AX455627	AX455627 Sequence
5	20	100.0	22	6	AR060840	AR060840 Sequence
6	20	100.0	22	6	AR070792	AR070792 Sequence
7	20	100.0	22	6	AX195274	AX195274 Sequence
8	20	100.0	48	6	AX377574	AX377574 Sequence
9	20	100.0	48	6	AX127457	AX127457 Sequence
10	20	100.0	49	6	AX127457	AX127457 Sequence
11	20	100.0	49	6	AX127457	AX127457 Sequence
12	20	100.0	49	6	AX127457	AX127457 Sequence
13	20	100.0	49	6	AX127457	AX127457 Sequence
14	20	100.0	49	6	AX127457	AX127457 Sequence
15	19	95.0	21	6	AX104633	AX104633 Sequence
16	19	95.0	21	6	AX104634	AX104634 Sequence
17	19	95.0	21	6	AX355087	AX355087 Sequence
18	19	95.0	21	6	AX355201	AX355201 Sequence
19	19	95.0	46	6	I72381	I72381 Sequence 12
20	18.4	92.0	22	6	E07877	E07877 Synthetic n
21	18	90.0	46	6	I72381	I72381 Sequence 13
22	18	90.0	46	6	I72382	I72382 Sequence 12
23	18	90.0	46	6	I72382	I72382 Sequence 13
24	16.8	84.0	49	6	AX377573	AX377573 Sequence
25	15.8	79.0	31	6	AR091898	AR091898 Sequence
26	15.8	79.0	31	6	E64477	E64477 Sugar-respo
27	15.8	79.0	31	6	I77207	I77207 Sequence 4
28	15.8	79.0	66	6	AX207285	AX207285 Sequence
29	15.2	76.0	80	6	AX002555	AX002555 Sequence
30	15.2	76.0	80	6	E27453	E27453 cdc25B Gene
31	14.8	74.0	27	6	AX182197	AX182197 Sequence
32	14.8	74.0	27	6	AX382003	AX382003 Sequence
33	14.4	72.0	58	6	AR034640	AR034640 Sequence
34	14.4	72.0	58	6	I70120	I70120 Sequence 26
35	14.4	72.0	60	6	AR034639	AR034639 Sequence
36	14.4	72.0	60	6	I70119	I70119 Sequence 25
37	13.6	68.0	33	6	AX363257	AX363257 Sequence
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39	13.6	68.0	65	6	AR121168	AR121168 Sequence
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42	13.2	66.0	39	6	AX006290	AX006290 Sequence
43	13.2	66.0	51	6	AX204497	AX204497 Sequence
44	13.2	66.0	81	14	AF221250	AF221250 Hepatitis
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# ALIGNMENTS

RESULT 1

A89791

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

A89791  
Sequence 13 from Patent WO9832462.  
A89791  
A89791.1 GI:6738305

20 bp

DNA

linear

PAT 22-JAN-2000

unidentified.

unclassified.

1 (bases 1 to 20)

Lipford, G.B. and Heeg, K.

PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND

OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION

Patent: WO 9832462-A 13 30-JUL-1998;

Pred. No. is the number of results predicted by chance to have a

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DEFINITION		Sequence 13 from Patent EP0855184.			
ACCESSION		A90878			
VERSION		A90878.1 GI:6739281			
KEYWORDS		unidentified.			
SOURCE		unidentified			
ORGANISM		unidentified			
REFERENCE		1 (bases 1 to 20)			
AUTHORS		Heeg, K.P. and Lipford, G.B.			
TITLE		Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination			
JOURNAL		Patent: EP 0855184-A 13 29-JUL-1998;			
FEATURES		HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)			
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LOCUS		AX023404		PAT 15-SEP-2000	
DEFINITION		Sequence 19 from Patent WO0014217.			
ACCESSION		AX023404			
VERSION		AX023404.1 GI:10183804			
KEYWORDS		synthetic construct.			
SOURCE		synthetic construct			
ORGANISM		artificial sequences.			
REFERENCE		1 (bases 1 to 20)			
AUTHORS		Lipford, G.B., Heeg, K. and Wagner, H.			
TITLE		G-motif oligonucleotides and uses thereof			
JOURNAL		Patent: WO 0014217-A 19 16-MAR-2000;			
FEATURES		LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);			
source		CPG IMMUNOPHARMACEUTICALS GMBH (DE)			
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/organism="synthetic construct"					
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/note="synthetic, no natural origin"					
BASE COUNT		2 a 5 c 11 g			
ORIGIN					

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)		Lipford, G. B. and Heeg, K. and Wagner, H.		PAT 22-JAN-2000	
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Best Local Similarity		100.0%; Pred. No. 3.3e+02;			
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Db		1 TCGATCGGGCGGGCGGAGC 20			
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LOCUS		A90878		PAT 22-JAN-2000	
DEFINITION		Sequence 13 from Patent EP0855184.			
ACCESSION		A90878			
VERSION		A90878.1 GI:6739281			
KEYWORDS		unidentified.			
SOURCE		unidentified			
ORGANISM		unidentified			
REFERENCE		1 (bases 1 to 20)			
AUTHORS		Heeg, K.P. and Lipford, G.B.			
TITLE		Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination			
JOURNAL		Patent: EP 0855184-A 13 29-JUL-1998;			
FEATURES		HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)			
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RESULT 3					
LOCUS		AX023404		PAT 15-SEP-2000	
DEFINITION		Sequence 19 from Patent WO0014217.			
ACCESSION		AX023404			
VERSION		AX023404.1 GI:10183804			
KEYWORDS		synthetic construct.			
SOURCE		synthetic construct			
ORGANISM		artificial sequences.			
REFERENCE		1 (bases 1 to 20)			
AUTHORS		Lipford, G.B., Heeg, K. and Wagner, H.			
TITLE		G-motif oligonucleotides and uses thereof			
JOURNAL		Patent: WO 0014217-A 19 16-MAR-2000;			

AR070792 LOCUS 22 bp DNA linear PAT 18-FEB-2000  
DEFINITION Sequence 12 from patent US 5908762.  
ACCESSION AR070792  
VERSION AR070792.1 GI:7221680  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 22)  
AUTHORS Ono, S., Jeremy, and Strominger, J. L.  
TITLE Transcription factor regulating MHC expression CDNA and genomic clones encoding same and retroviral expression constructs thereof  
JOURNAL Patent: US 5908762-A 12 01-JUN-1999;  
FEATURES Location/Qualifiers  
source 1..22  
BASE COUNT 3 a 5 c 11 g 3 t  
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Query Match 100.0%; Score 20; DB 6; Length 22;  
Best Local Similarity 100.0%; Pred. No. 3.2e+02;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 TCGATCGGGCGGGCGGAGC 20  
Db 3 TCGATCGGGCGGGCGGAGC 22  
RESULT 7  
LOCUS AX195274 22 bp DNA linear PAT 28-AUG-2001  
DEFINITION Sequence 10 from Patent WO0151671.  
ACCESSION AX195274  
VERSION AX195274.1 GI:15385825  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 22)  
AUTHORS McCarthy, J. and Cordell, B.  
TITLE Methods for identifying inhibitors of neuronal degeneration  
JOURNAL Patent: WO 0151671-A 10 19-JUL-2001;  
FEATURES Scios Inc. (US)  
source Location/Qualifiers  
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/note="Synthetic Oligonucleotide"  
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Best Local Similarity 100.0%; Pred. No. 3.2e+02;  
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Db 3 TCGATCGGGCGGGCGGAGC 22  
RESULT 8  
LOCUS AX377574 48 bp DNA linear PAT 18-MAR-2002  
DEFINITION Sequence 51 from Patent WO0212553.  
ACCESSION AX377574  
VERSION AX377574.1 GI:19573760  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 49)  
AUTHORS Muth, J., and Windhab, N.  
TITLE Double-strand nucleic acid probes and the use thereof  
JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;  
FEATURES Aventis Research & Technologies GmbH & Co KG (DE)  
source Location/Qualifiers  
1..49

AR070792 LOCUS 22 bp DNA linear PAT 18-FEB-2000  
DEFINITION Sequence 12 from patent US 5908762.  
ACCESSION AR070792  
VERSION AR070792.1 GI:7221680  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 22)  
AUTHORS Ono, S., Jeremy, and Strominger, J. L.  
TITLE Transcription factor regulating MHC expression CDNA and genomic clones encoding same and retroviral expression constructs thereof  
JOURNAL Patent: US 5908762-A 12 01-JUN-1999;  
FEATURES Location/Qualifiers  
source 1..22  
BASE COUNT 3 a 5 c 11 g 3 t  
ORIGIN  
Query Match 100.0%; Score 20; DB 6; Length 22;  
Best Local Similarity 100.0%; Pred. No. 3.2e+02;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 TCGATCGGGCGGGCGGAGC 20  
Db 3 TCGATCGGGCGGGCGGAGC 22  
RESULT 7  
LOCUS AX195274 22 bp DNA linear PAT 28-AUG-2001  
DEFINITION Sequence 10 from Patent WO0151671.  
ACCESSION AX195274  
VERSION AX195274.1 GI:15385825  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 22)  
AUTHORS McCarthy, J. and Cordell, B.  
TITLE Methods for identifying inhibitors of neuronal degeneration  
JOURNAL Patent: WO 0151671-A 10 19-JUL-2001;  
FEATURES Scios Inc. (US)  
source Location/Qualifiers  
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/note="Synthetic Oligonucleotide"  
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ORIGIN  
Query Match 100.0%; Score 20; DB 6; Length 22;  
Best Local Similarity 100.0%; Pred. No. 3.2e+02;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 TCGATCGGGCGGGCGGAGC 20  
Db 3 TCGATCGGGCGGGCGGAGC 22  
RESULT 8  
LOCUS AX377574 48 bp DNA linear PAT 18-MAR-2002  
DEFINITION Sequence 51 from Patent WO0212553.  
ACCESSION AX377574  
VERSION AX377574.1 GI:19573760  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 49)  
AUTHORS Muth, J., and Windhab, N.  
TITLE Double-strand nucleic acid probes and the use thereof  
JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;  
FEATURES Aventis Research & Technologies GmbH & Co KG (DE)  
source Location/Qualifiers  
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Muth, J.  
Method for detecting mutations in nucleotide sequences  
Patent: WO 0212553-A 51 14-FEB-2002;  
Nanogen Recognomics GmbH (DE)  
Location/Qualifiers  
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ACCESSION AX377574  
VERSION AX377574.1 GI:19573760  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 48)  
AUTHORS Kappel, A., Polakowski, T., Pignot, M., Windhab, N., Behrens, H., and Muth, J.  
TITLE Method for detecting mutations in nucleotide sequences  
JOURNAL Patent: WO 0212553-A 51 14-FEB-2002;  
Nanogen Recognomics GmbH (DE)  
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ACCESSION AX127457  
VERSION AX127457.1 GI:14134020  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 49)  
AUTHORS Muth, J., and Windhab, N.  
TITLE Double-strand nucleic acid probes and the use thereof  
JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;  
FEATURES Aventis Research & Technologies GmbH & Co KG (DE)  
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ACCESSION AX127457
VERSION AX127457.1 GI:14134020
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 49)
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 1 03-MAY-2001;
Aventis Research & Technologies GmbH & Co KG (DE)
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DEFINITION Sequence 3 from Patent WO0131057.
ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 49)
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 3 03-MAY-2001;
Aventis Research & Technologies GmbH & Co KG (DE)
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ACCESSION AX127459
VERSION AX127459.1 GI:14134028
KEYWORDS
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ORGANISM synthetic construct
artificial sequences.
REFERENCE 1 (bases 1 to 49)
AUTHORS Muth,J. and Windhab,N.
TITLE Double-strand nucleic acid probes and the use thereof
JOURNAL Patent: WO 0131057-A 3 03-MAY-2001;
Aventis Research & Technologies GmbH & Co KG (DE)
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Db 47 TCGATCGGGCGGGCGGCGAGC 28

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DEFINITION Sequence 50 from Patent WO0212553.
ACCESSION AX377573
VERSION AX377573.1 GI:19573759
KEYWORDS
SOURCE synthetic construct.
ORGANISM synthetic construct
artificial sequences.
REFERENCE 1
AUTHORS Kappel,A., Polakowski,T., Pignot,M., Windhab,N., Behrensdoerf,H. and
Muth,J.
TITLE Method for detecting mutations in nucleotide sequences
JOURNAL Patent: WO 0212553-A 50 14-FEB-2002;
Nanogen Recognomics GmbH (DE)
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Location/Qualifiers
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GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:05:40 ; Search time 22.2464 Seconds  
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Scoring table: IDENTITY\_NUC  
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Searched: 441362 seqs, 15338381 residues

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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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4	20	100.0	21	3	US-08-764-528-2
5	20	100.0	21	3	US-08-872-859-2
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15	13.6	68.0	26	1	US-07-791-213D-68
16	13.6	68.0	26	1	US-08-293-150A-68
17	13.4	67.0	41	2	US-08-818-604-8
18	12.8	64.0	18	3	US-09-289-377-17
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23	12.8	64.0	44	5	PCT-US95-02945-17
24	12.6	63.0	20	4	US-09-326-186B-171
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31	12.4	62.0	37	1	US-08-203-534-2	Sequence 2, Appl
32	12.4	62.0	41	2	US-08-350-260A-586	Sequence 586, App
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ALIGNMENTS

RESULT 1  
US-08-507-598-2  
; Sequence 2, Application US/08507598  
; Patent No. 5834188  
; GENERAL INFORMATION:  
; APPLICANT: HARADA, SHUN-ICHI  
; APPLICANT: SAMPATH, T. K.  
; APPLICANT: RODAN, GIDEON A.  
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING  
; TITLE OF INVENTION: MORPHOGEN ANALOGS  
; NUMBER OF SEQUENCES: 4  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &  
; ADDRESS: THIBEAULT  
; STREET: 53 STATE STREET  
; CITY: BOSTON  
; STATE: MA  
; COUNTRY: USA  
; ZIP: 02109  
; COMPUTER READABLE FORM:  
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; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/507,598  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: PITCHER, EDMUND R.  
; REGISTRATION NUMBER: 27,829  
; REFERENCE/DOCKET NUMBER: CRP-107  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (617)-248-7000  
; TELEFAX: (617)-248-7100  
; INFORMATION FOR SEQ ID NO: 2:  
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; LENGTH: 21 base pairs  
; TYPE: nucleic acid  
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## RESULT 2

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; Patent No. 5932716  
; GENERAL INFORMATION:  
; APPLICANT: SAMPATH, T. K.  
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING  
; MORPHOGEN ANALOGS  
; NUMBER OF SEQUENCES: 4  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &  
; ADDRESSEE: THIBEAULT  
; STREET: 53 STATE STREET  
; CITY: BOSTON  
; STATE: MA  
; COUNTRY: USA  
; ZIP: 02109

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; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/507,750  
; FILING DATE:  
; CLASSIFICATION: 424  
; ATTORNEY/AGENT INFORMATION:  
; NAME: FITCHER, EDMUND R.  
; REGISTRATION NUMBER: 27,829  
; REFERENCE/DOCKET NUMBER: CRP-116  
; TELEPHONE: (617)-248-7000  
; TELEFAX: (617)-248-7100

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; MOLECULE TYPE: DNA (genomic)  
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; Sequence 2, Application US/08764522A  
; Patent No. 6090544  
; GENERAL INFORMATION:  
; APPLICANT: HARADA, SHUN-ICHI  
; APPLICANT: RODAN, GIDEON A.  
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING  
; MORPHOGEN ANALOGS  
; NUMBER OF SEQUENCES: 10  
; FILING DATE:  
; CLASSIFICATION: 424  
; ATTORNEY/AGENT INFORMATION:  
; NAME: VITO, CHRISTINE C.  
; REGISTRATION NUMBER: 39,061  
; REFERENCE/DOCKET NUMBER: CRP-126  
; TELEPHONE: (617)-248-7000  
; TELEFAX: (617)-248-7100

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; APPLICANT: HARADA, SHUN-ICHI  
; APPLICANT: RODAN, GIDEON A.  
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING  
; MORPHOGEN ANALOGS  
; NUMBER OF SEQUENCES: 10  
; FILING DATE:  
; CLASSIFICATION: 424  
; ATTORNEY/AGENT INFORMATION:  
; NAME: VITO, CHRISTINE C.  
; REGISTRATION NUMBER: 39,061

CORRESPONDENCE ADDRESS:  
; ADDRESSEE: PATENT ADMINISTRATOR, CREATIVE BIOMOLECULES  
; STREET: 45 SOUTH STREET  
; CITY: HOPKINTON  
; STATE: MA  
; COUNTRY: USA  
; ZIP: 01748

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; APPLICATION NUMBER: US/08/764,522A  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: VITO, CHRISTINE C.  
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; REFERENCE/DOCKET NUMBER: CRP-126  
; TELEPHONE: (617)-248-7000  
; TELEFAX: (617)-248-7100

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; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
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RESULT 4  
US-08-764-528-2

; Sequence 2, Application US/08764528  
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; APPLICANT: SAMPATH, K. T.  
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING  
; MORPHOGEN ANALOGS  
; NUMBER OF SEQUENCES: 8  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: PATENT ADMINISTRATOR, CREATIVE BIOMOLECULES  
; STREET: 45 SOUTH STREET  
; CITY: HOPKINTON  
; STATE: MA  
; COUNTRY: USA  
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; FILING DATE:  
; CLASSIFICATION: 424  
; ATTORNEY/AGENT INFORMATION:  
; NAME: VITO, CHRISTINE C.  
; REGISTRATION NUMBER: 39,061

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; REFERENCE/DOCKET NUMBER: CRP-127
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; TELEFAX: (617)-248-7100
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Db 2 GCTTGATGACTCAGCGGAA 21

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; Patent No. 6110460
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; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR IDENTIFYING
; TITLE OF INVENTION: MORPHOGEN ANALOGS
; NUMBER OF SEQUENCES: 4
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: PATENT ADMINISTRATOR, TESTA, HURWITZ &
; ADDRESSEE: THIBEAULT
; STREET: 53 STATE STREET
; CITY: BOSTON
; STATE: MA
; COUNTRY: USA
; ZIP: 02109
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; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/507,750
; FILING DATE: 26-JUL-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: PITCHER, EDMUND R.
; REGISTRATION NUMBER: 27,829
; REFERENCE/DOCKET NUMBER: CRP-116
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (617)-248-7000
; TELEFAX: (617)-248-7100
; INFORMATION FOR SEQ ID NO: 2:
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US-08-872-859-2

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Best Local Similarity 100.0%; Pred. No. 0.058;
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QY 1 GCTTGATGACTCAGCGGAA 20
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RESULT 6
US-08-283-591-15
; Sequence 15, Application US/08283591
; Patent No. 5629152
; GENERAL INFORMATION:
; APPLICANT: Ravikumar, Vasulunga
; TITLE OF INVENTION: NOVEL TRISUBSTITUTED -LACTAMS AND
; TITLE OF INVENTION: OLIGO -LACTAMAMIDES
; NUMBER OF SEQUENCES: 28
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Woodcock Washburn Kurtz Mackiewicz and
; ADDRESSEE: No. 5629152rls
; STREET: One Liberty Place - 46th Floor
; CITY: Philadelphia
; STATE: PA
; COUNTRY: U.S.A.
; ZIP: 19103
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/283,591
; FILING DATE: N/A
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME:
; REGISTRATION NUMBER:
; REFERENCE/DOCKET NUMBER:
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 215-568-3100
; TELEFAX: 215-568-3439
; INFORMATION FOR SEQ ID NO: 15:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 21
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: unknown
US-08-283-591-15

Query Match 92.0%; Score 18.4; DB 1; Length 21;
Best Local Similarity 95.0%; Pred. No. 0.42;
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20
Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 7
US-08-210-880B-3
; Sequence 3, Application US/08210880B
; Patent No. 5641486
; GENERAL INFORMATION:
; APPLICANT: HINRICHS, STEVEN H.
; APPLICANT: ORTEN, DANA J.
; TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING
; TITLE OF INVENTION: CREB/ATF1 FAMILY DETERMINED TRANSCRIPTION

```

NUMBER OF SEQUENCES: 11  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: HENDERSON & STURM  
STREET: 1125 S. 103RD ST., #330  
CITY: OMAHA  
STATE: NE  
COUNTRY: US  
ZIP: 68124  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/210.880B  
FILING DATE: 18-MAR-1994  
CLASSIFICATION: 424  
ATTORNEY/AGENT INFORMATION:  
NAME: JONDLE, ROBERT J.  
REGISTRATION NUMBER: 33,915  
REFERENCE/DOCKET NUMBER: 63066  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 402-398-9000  
TELEFAX: 402-398-9005  
INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 21 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: unknown  
MOLECULE TYPE: DNA (genomic)  
US-08-210-880B-3

Query Match 92.0%; Score 18.4; DB 1; Length 21;  
Best Local Similarity 95.0%; Pred. No. 0.42;  
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GCTTGATGACTCAGCGGAA 20  
||||| |||||||  
Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 8  
US-08-632-275-1/c  
Sequence 1, Application US/08632275  
Patent No. 5840277  
GENERAL INFORMATION:  
APPLICANT: Ghio, Andrew J.  
ADDRESSEE: Kennedy, Thomas P.  
TITLE OF INVENTION: Treatment of Chronic Pulmonary  
TITLE OF INVENTION: Inflammation  
NUMBER OF SEQUENCES: 4  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Bell, Seltzer, Park & Gibson  
STREET: 1211 East Morehead Street  
CITY: Charlotte  
STATE: No. 5840277th Carolina  
COUNTRY: USA  
ZIP: 28234  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/632.275  
FILING DATE: 15-APR-1996  
CLASSIFICATION: 514  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/413.699  
FILING DATE: 30-MAR-1995  
CLASSIFICATION: 514  
ATTORNEY/AGENT INFORMATION:

NAME: Lipscomb, Ernest B.  
REGISTRATION NUMBER: 24,733  
REFERENCE/DOCKET NUMBER: 8751-5-1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 704-331-6000  
TELEFAX: 704-334-2014  
INFORMATION FOR SEQ ID NO: 1:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 21 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
FRAGMENT TYPE: linear  
US-08-632-275-1  
Query Match 92.0%; Score 18.4; DB 2; Length 21;  
Best Local Similarity 95.0%; Pred. No. 0.42;  
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 1 GCTTGATGACTCAGCGGAA 20  
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Db 20 GCTTGATGACTCAGCGGAA 1

RESULT 9  
US-08-771-411-3  
Sequence 3, Application US/08771411  
Patent No. 5844096  
GENERAL INFORMATION:  
APPLICANT: HINRICHS, STEVEN H.  
APPLICANT: ORTEN, DANA J.  
TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING  
TITLE OF INVENTION: CREB/ATF1 FAMILY DETERMINED TRANSCRIPTION  
NUMBER OF SEQUENCES: 11  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: HENDERSON & STURM  
STREET: 1125 S. 103RD ST., #330  
CITY: OMAHA  
STATE: NE  
COUNTRY: US  
ZIP: 68124  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/771.411  
FILING DATE: 20-DEC-1996  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/210.880  
FILING DATE: 18-MAR-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: JONDLE, ROBERT J.  
REGISTRATION NUMBER: 33,915  
REFERENCE/DOCKET NUMBER: 63066  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 402-398-9000  
TELEFAX: 402-398-9005  
INFORMATION FOR SEQ ID NO: 3:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 21 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: unknown  
MOLECULE TYPE: DNA (genomic)  
US-08-771-411-3

Query Match 92.0%; Score 18.4; DB 2; Length 21;  
Best Local Similarity 95.0%; Pred. No. 0.42;  
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 10

US-09-097-929-1/c  
 ; Sequence 1, Application US/09097929  
 ; Patent No. 6024940  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Ghio, Andrew J.  
 ; APPLICANT: Kennedy, Thomas P.  
 ; TITLE OF INVENTION: Treatment of Chronic Pulmonary  
 ; TITLE OF INVENTION: Inflammation  
 ; NUMBER OF SEQUENCES: 4  
 ; CORRESPONDENCE ADDRESS:  
 ; ADDRESSEE: Bell, Seltzer, Park & Gibson  
 ; STREET: 1211 East Morehead Street  
 ; CITY: Charlotte  
 ; STATE: No. 6024940th Carolina  
 ; COUNTRY: USA  
 ; ZIP: 28234  
 ; COMPUTER READABLE FORM:  
 ; MEDIUM TYPE: Floppy disk  
 ; COMPUTER: IBM PC compatible  
 ; OPERATING SYSTEM: PC-DOS/MS-DOS  
 ; SOFTWARE: PatentIn Release #1.0, Version #1.30  
 ; CURRENT APPLICATION DATA:  
 ; APPLICATION NUMBER: US/09/097,929  
 ; FILING DATE:  
 ; CLASSIFICATION:  
 ; PRIOR APPLICATION DATA:  
 ; APPLICATION NUMBER: 08/632,275  
 ; FILING DATE:  
 ; CLASSIFICATION:  
 ; ATTORNEY/AGENT INFORMATION:  
 ; NAME: Lipscomb, Ernest B.  
 ; REGISTRATION NUMBER: 24,733  
 ; REFERENCE/DOCKET NUMBER: 8751-5-1  
 ; TELECOMMUNICATION INFORMATION:  
 ; TELEPHONE: 704-331-6000  
 ; TELEFAX: 704-334-2014  
 ; INFORMATION FOR SEQ ID NO: 1:  
 ; SEQUENCE CHARACTERISTICS:  
 ; LENGTH: 21 base pairs  
 ; TYPE: nucleic acid  
 ; STRANDEDNESS: single  
 ; TOPOLOGY: linear  
 ; FRAGMENT TYPE: linear  
 ; US-09-097-929-1

Query Match 92.0%; Score 18.4; DB 3; Length 21;  
 Best Local Similarity 95.0%; Pred. No. 0.42; Mismatches 0; Gaps 0;  
 Matches 19; Conservative 0; Indels 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 Db 20 GCTTGATGACTCAGCGGAA 1

RESULT 11

US-09-021-247-8  
 ; Sequence 8, Application US/09021247  
 ; Patent No. 6225444  
 ; GENERAL INFORMATION:  
 ; APPLICANT: Shastoua, Victor E.  
 ; TITLE OF INVENTION: NEUROPROTECTIVE PEPTIDES AND USES THEREOF  
 ; NUMBER OF SEQUENCES: 19  
 ; CORRESPONDENCE ADDRESS:  
 ; ADDRESSEE: Wolf, Greenfield & Sacks, P.C.  
 ; STREET: 600 Atlantic Avenue  
 ; CITY: Boston  
 ; STATE: MA

; COUNTRY: USA  
 ; ZIP: 02210  
 ; COMPUTER READABLE FORM:  
 ; MEDIUM TYPE: Floppy disk  
 ; COMPUTER: IBM PC compatible  
 ; OPERATING SYSTEM: PC-DOS/MS-DOS  
 ; SOFTWARE: PatentIn Release #1.0, Version #1.25  
 ; CURRENT APPLICATION DATA:  
 ; APPLICATION NUMBER: US/09/021,247  
 ; FILING DATE:  
 ; CLASSIFICATION:  
 ; ATTORNEY/AGENT INFORMATION:  
 ; NAME: Van Amsterdam, John R.  
 ; REGISTRATION NUMBER: 40,212  
 ; REFERENCE/DOCKET NUMBER: N0260/7023  
 ; TELECOMMUNICATION INFORMATION:  
 ; TELEPHONE: 617-720-3500  
 ; TELEFAX: 617-720-2441  
 ; INFORMATION FOR SEQ ID NO: 8:  
 ; SEQUENCE CHARACTERISTICS:  
 ; LENGTH: 21 nucleotides  
 ; TYPE: nucleic acid  
 ; STRANDEDNESS: single  
 ; TOPOLOGY: linear  
 ; MOLECULE TYPE: oligonucleotide  
 ; HYPOTHETICAL: NO  
 ; US-09-021-247-8

Query Match 92.0%; Score 18.4; DB 4; Length 21;  
 Best Local Similarity 95.0%; Pred. No. 0.42; Mismatches 0; Gaps 0;  
 Matches 19; Conservative 0; Indels 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 12

US-08-088-661F-8  
 ; Sequence 8, Application US/08088661F  
 ; Patent No. 6228982  
 ; GENERAL INFORMATION:  
 ; APPLICANT: No. 6228982den, Bengt  
 ; APPLICANT: Wittung, Pernilla  
 ; APPLICANT: Buchardt, Ole  
 ; APPLICANT: Egholm, Michael  
 ; APPLICANT: Nielsen, Peter E.  
 ; APPLICANT: Berg, Rolf  
 ; TITLE OF INVENTION: Double-Stranded Peptide Nucleic Acids  
 ; FILE REFERENCE: ISIS1108  
 ; CURRENT APPLICATION NUMBER: US/08/088,661F  
 ; CURRENT FILING DATE: 1993-07-02  
 ; PRIOR APPLICATION NUMBER: 08/054,363  
 ; PRIOR FILING DATE: 1993-04-26  
 ; PRIOR APPLICATION NUMBER: PCT/EP92/01219  
 ; PRIOR FILING DATE: 1992-05-19  
 ; NUMBER OF SEQ ID NOS: 42  
 ; SOFTWARE: PatentIn Ver. 2.1  
 ; SEQ ID NO 8  
 ; LENGTH: 21  
 ; TYPE: DNA  
 ; ORGANISM: Artificial Sequence  
 ; FEATURE:  
 ; OTHER INFORMATION: Description of Artificial Sequence: No. 6228982el Sequence  
 ; US-08-088-661F-8

Query Match 92.0%; Score 18.4; DB 4; Length 21;  
 Best Local Similarity 95.0%; Pred. No. 0.42; Mismatches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
 Db 2 GCTTGATGACTCAGCGGAA 21

RESULT 13  
US-08-088-661F-42  
; Sequence 42, Application US/08088661F  
; Patent No. 6228982  
; GENERAL INFORMATION:  
; APPLICANT: No. 6228982den, Benget  
; APPLICANT: Wittung, Pernilla  
; APPLICANT: Buchardt, Ole  
; APPLICANT: Egholm, Michael  
; APPLICANT: Nielsen, Peter E.  
; APPLICANT: Berg, Rolf  
; TITLE OF INVENTION: Double-Stranded Peptide Nucleic Acids  
; FILE REFERENCE: IS11108  
; CURRENT APPLICATION NUMBER: US/08/088,661F  
; CURRENT FILING DATE: 1993-07-02  
; PRIOR APPLICATION NUMBER: 08/054,363  
; PRIOR FILING DATE: 1993-04-26  
; PRIOR APPLICATION NUMBER: PCT/EP92/01219  
; PRIOR FILING DATE: 1992-05-19  
; NUMBER OF SEQ ID NOS: 42  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 42  
; LENGTH: 21  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: Description of Artificial Sequence: No. 6228982el Sequence  
US-08-088-661F-42

Query Match 92.0%; Score 18.4; DB 4; Length 21;  
Best Local Similarity 95.0%; Pred. No. 0.42;  
Matches 19; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20  
||||| ||||||| |||||  
Db 2 GCTTGGTGACTCAGCGGAA 21

RESULT 14  
US-08-203-198-1  
; Sequence 1, Application US/08203198  
; Patent No. 5512462  
; GENERAL INFORMATION:  
; APPLICANT: Cheng, Suzanne  
; TITLE OF INVENTION: Methods and Reagents for the Polymerase  
; NUMBER OF SEQUENCES: 32  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Hoffmann-La Roche Inc.  
; STREET: 340 Kingsland Street  
; CITY: Nutley  
; STATE: NJ  
; COUNTRY: U.S.A.  
; ZIP: 07110  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/08/203,198  
; FILING DATE:  
; CLASSIFICATION: 435  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Petry Ph.D., Douglas A.  
; REGISTRATION NUMBER: 35,321  
; REFERENCE/DOCKET NUMBER: 8694  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (510) 814-2974  
; TELEFAX: (510) 814-2977  
; INFORMATION FOR SEQ ID NO: 1:

; SEQUENCE CHARACTERISTICS:  
; LENGTH: 21 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
US-08-203-198-1

Query Match 69.0%; Score 13.8; DB 1; Length 21;  
Best Local Similarity 88.2%; Pred. No. 1.2e+02;  
Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCG 17  
||||| ||||||| |||||  
Db 4 GCTTTATGACTCTGCG 20

## RESULT 15

US-07-791-213D-68/C  
; Sequence 68, Application US/07791213D  
; Patent No. 5409895  
; GENERAL INFORMATION:  
; APPLICANT: MORISHITA, Hideaki  
; APPLICANT: KANAMORI, Toshinori  
; APPLICANT: NOBUHARA, Masahiro  
; TITLE OF INVENTION: POLYPEPTIDE, DNA FRAGMENT ENCODING THE  
; TITLE OF INVENTION: SAME AND PROCESS FOR PRODUCING THE SAME, AND ENZYME  
; TITLE OF INVENTION: INHIBITION PROCESS, DRUG COMPOSITION AND METHODS OF  
; TITLE OF INVENTION: TREATING USING THE SAME  
; NUMBER OF SEQUENCES: 108  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Burns, Doane, Swecker & Mathis  
; STREET: P.O. Box 1404  
; CITY: Alexandria  
; STATE: Virginia  
; COUNTRY: United States  
; ZIP: 22313-1404  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.25  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/07/791,213D  
; FILING DATE: 13-NOV-1991  
; CLASSIFICATION: 435  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: JP 2-306745  
; FILING DATE: 13-NOV-1990  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Meuth, Donna M  
; REGISTRATION NUMBER: 36,607  
; REFERENCE/DOCKET NUMBER: 029650-032  
; TELECOMMUNICATION INFORMATION:  
; TELEPHONE: (703) 836-6620  
; TELEFAX: (703) 836-2021  
; INFORMATION FOR SEQ ID NO: 68:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 26 base pairs  
; TYPE: nucleic acid  
; STRANDEDNESS: single  
; TOPOLOGY: linear  
; MOLECULE TYPE: DNA (genomic)  
; FEATURE:  
; NAME/KEY: misc\_feature  
; LOCATION: complement (1..26)  
US-07-791-213D-68

Query Match 68.0%; Score 13.6; DB 1; Length 26;  
Best Local Similarity 80.0%; Pred. No. 1.6e+02;  
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 GCTTGATGACTCAGCGGAA 20

Db 26 GCTGGATCCCTCAGCCGAAA 7

Search completed: December 12, 2002, 01:41:48  
Job time : 24.2464 secs





GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds  
(without alignments)  
1829.698 Million cell updates/sec

Title: US-09-355-254F-14

Perfect score: 20  
Sequence: 1 agcggggcgagcgggggcg 20

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

GenEmbl.\*

- 1: gb\_ba.\*
- 2: gb\_hgt.\*
- 3: gb\_in.\*
- 4: gb\_om.\*
- 5: gb\_ov.\*
- 6: gb\_pat.\*
- 7: gb\_ph.\*
- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
- 12: gb\_sy.\*
- 13: gb\_un.\*
- 14: gb\_vi.\*
- 15: em\_ba.\*
- 16: em\_fun.\*
- 17: em\_hum.\*
- 18: em\_in.\*
- 19: em\_mu.\*
- 20: em\_om.\*
- 21: em\_or.\*
- 22: em\_ov.\*
- 23: em\_pat.\*
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- 27: em\_sts.\*
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- 29: em\_vi.\*
- 30: em\_hgt\_hum.\*
- 31: em\_hgt\_inv.\*
- 32: em\_hgt\_other.\*
- 33: em\_hgt\_mus.\*
- 34: em\_hgt\_pln.\*
- 35: em\_hgt\_rtd.\*
- 36: em\_hgt\_mam.\*
- 37: em\_hgt\_vrt.\*
- 38: em\_sy.\*
- 39: em\_htgo\_hum.\*
- 40: em\_htgo\_mus.\*
- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	20	100.0	20	6 A89793	A89793 Sequence 15
2	20	100.0	20	6 A90880	A90880 Sequence 15
3	20	100.0	20	6 AX023403	AX023403 Sequence
4	20	100.0	20	6 AX455554	AX455554 Sequence
5	20	100.0	30	6 AX472520	AX472520 Sequence
6	20	100.0	30	6 AX476834	AX476834 Sequence
7	20	100.0	30	6 AX476857	AX476857 Sequence
8	16.4	82.0	30	6 AX063474	AX063474 Sequence
9	16.4	82.0	30	6 I28154	I28154 Sequence 11
10	14.2	71.0	47	6 AX128389	AX128389 Sequence
11	14.2	71.0	50	6 AR032813	AR032813 Sequence
12	14.2	71.0	50	6 AR209477	AR209477 Sequence
13	14.2	71.0	50	6 I29553	I29553 Sequence 42
14	14.2	71.0	50	6 I91227	I91227 Sequence 42
15	14.2	71.0	51	6 AX158283	AX158283 Sequence
16	13.8	69.0	72	9 HSDNASDAE	Z22319 H.sapiens D
17	13.8	69.0	100	11 AF235063	AF235063 Mus muscu
18	13.6	68.0	30	6 AX472521	AX472521 Sequence
19	13.6	68.0	30	6 AX476835	AX476835 Sequence
20	13.6	68.0	40	6 I86249	I86249 Sequence 3
21	13.6	68.0	44	6 AX157237	AX157237 Sequence
22	13.6	68.0	50	10 MMU41966	U41966 Mus musculu
23	13.6	68.0	51	6 A42081	A42081 Sequence 24
24	13.6	68.0	98	6 AR017637	AR017637 Sequence
25	13.6	68.0	98	6 AR094814	AR094814 Sequence
26	13.6	68.0	98	6 AR165473	AR165473 Sequence
27	13.6	68.0	99	8 AY033466	AY033466 zea mays
28	13.6	68.0	100	6 I28243	I28243 Sequence 1
29	13.4	67.0	24	6 AR058202	AR058202 Sequence
30	13.4	67.0	24	6 AR152031	AR152031 Sequence
31	13.4	67.0	24	6 I50793	I50793 Sequence 24
32	13.4	67.0	30	6 AX167104	AX167104 Sequence
33	13.4	67.0	33	6 AX202261	AX202261 Sequence
34	13.2	66.0	19	6 AX129905	AX129905 Sequence
35	13.2	66.0	19	6 AX129926	AX129926 Sequence
36	13.2	66.0	19	6 AX202627	AX202627 Sequence
37	13.2	66.0	21	6 A88115	A88115 Sequence 26
38	13.2	66.0	21	6 A90082	A90082 Sequence 26
39	13.2	66.0	22	6 A33211	A33211 Synthetic H
40	13.2	66.0	47	6 AR154447	AR154447 Sequence
41	13.2	66.0	50	6 AX159936	AX159936 Sequence
42	13.2	66.0	51	6 AX159935	AX159935 Sequence
43	13.2	66.0	54	6 AR032427	AR032427 Sequence
44	13.2	66.0	54	6 AR032648	AR032648 Sequence
45	13.2	66.0	54	6 AR209091	AR209091 Sequence

ALIGNMENTS

RESULT 1  
A89793  
LOCUS A89793  
DEFINITION Sequence 15 from Patent WO9832462.  
ACCESSION A89793  
VERSION A89793.1 GI:6738307  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unclassified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Lipford, G. B. and Heeg, K.  
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
JOURNAL Patent: WO 9832462-A 15 30-JUL-1998;

linear PAT 22-JAN-2000

## LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)

FEATURES  
Source

LOCUS AX023403 20 bp DNA linear PAT 15-SEP-2000  
 DEFINITION Sequence 18 from Patent WO0014217.  
 ACCESSION AX023403  
 VERSION AX023403.1 GI:10183803  
 KEYWORDS synthetic construct.  
 SOURCE artificial sequences.  
 REFERENCE 1 (bases 1 to 20)  
 AUTHORS Lipford, G.B., Heeg, K. and Wagner, H.  
 TITLE G-motif oligonucleotides and uses thereof  
 JOURNAL Patent: WO 0014217-A 18 16-MAR-2000;  
 LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);  
 CPG IMMUNOPHARMACEUTICALS GMBH (DE)

Query Match 100.0%; Score 20; DB 6; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 2.2e+03;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20  
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 Db 1 AGCGGGGCGAGCGGGGCG 20

## RESULT 2

LOCUS A90880 20 bp DNA linear PAT 22-JAN-2000  
 DEFINITION Sequence 15 from Patent EP0855184.  
 ACCESSION A90880  
 VERSION A90880.1 GI:6739295  
 KEYWORDS unclassified.  
 SOURCE unclassified.  
 ORGANISM unclassified.  
 REFERENCE 1 (bases 1 to 20)  
 AUTHORS Heeg, K.P. and Lipford, G.B.  
 TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination  
 JOURNAL Patent: EP 0855184-A 15 29-JUL-1998;  
 HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)

FEATURES  
Source

LOCUS AX023403 20 bp DNA linear PAT 15-SEP-2000  
 DEFINITION Sequence 18 from Patent WO0014217.  
 ACCESSION AX023403  
 VERSION AX023403.1 GI:10183803  
 KEYWORDS synthetic construct.  
 SOURCE artificial sequences.  
 REFERENCE 1 (bases 1 to 20)  
 AUTHORS Lipford, G.B., Heeg, K. and Wagner, H.  
 TITLE G-motif oligonucleotides and uses thereof  
 JOURNAL Patent: WO 0014217-A 18 16-MAR-2000;  
 LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);  
 CPG IMMUNOPHARMACEUTICALS GMBH (DE)

Query Match 100.0%; Score 20; DB 6; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 2.2e+03;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20  
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 Db 1 AGCGGGGCGAGCGGGGCG 20

## RESULT 3

LOCUS AX023403 20 bp DNA linear PAT 15-SEP-2000  
 DEFINITION Sequence 18 from Patent WO0014217.  
 ACCESSION AX023403  
 VERSION AX023403.1 GI:10183803  
 KEYWORDS synthetic construct.  
 SOURCE artificial sequences.  
 REFERENCE 1 (bases 1 to 20)  
 AUTHORS Lipford, G.B., Heeg, K. and Wagner, H.  
 TITLE G-motif oligonucleotides and uses thereof  
 JOURNAL Patent: WO 0014217-A 18 16-MAR-2000;  
 LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);  
 CPG IMMUNOPHARMACEUTICALS GMBH (DE)

FEATURES  
Source

LOCUS AX023403 20 bp DNA linear PAT 15-SEP-2000  
 DEFINITION Sequence 18 from Patent WO0014217.  
 ACCESSION AX023403  
 VERSION AX023403.1 GI:10183803  
 KEYWORDS synthetic construct.  
 SOURCE artificial sequences.  
 REFERENCE 1 (bases 1 to 20)  
 AUTHORS Lipford, G.B., Heeg, K. and Wagner, H.  
 TITLE G-motif oligonucleotides and uses thereof  
 JOURNAL Patent: WO 0014217-A 18 16-MAR-2000;  
 LIPFORD GRAYSON B (DE); HEEG KLAUS (DE); WAGNER HERMANN (DE);  
 CPG IMMUNOPHARMACEUTICALS GMBH (DE)

Query Match 100.0%; Score 20; DB 6; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 2.2e+03;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20  
 |||||  
 Db 1 AGCGGGGCGAGCGGGGCG 20

Query Match 100.0%; Score 20; DB 6; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 2.2e+03;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AGCGGGGCGAGCGGGGCG 20  
 |||||  
 Db 1 AGCGGGGCGAGCGGGGCG 20

## RESULT 4

LOCUS AX455554 20 bp DNA linear PAT 06-JUL-2002  
 DEFINITION Sequence 31 from Patent WO0222809.  
 ACCESSION AX455554  
 VERSION AX455554.1 GI:21714622  
 KEYWORDS synthetic construct.  
 SOURCE synthetic construct.  
 ORGANISM artificial sequences.  
 REFERENCE 1  
 AUTHORS Bauer, S., Lipford, G. and Wagner, H.  
 TITLE Process for high throughput screening of cpg-based immuno-agonist/antagonist  
 JOURNAL Patent: WO 0222809-A 31 21-MAR-2002;  
 Coley Pharmaceutical GmbH (DE)

FEATURES  
Location/Qualifiers

LOCUS AX455554 20 bp DNA linear PAT 06-JUL-2002  
 DEFINITION Sequence 31 from Patent WO0222809.  
 ACCESSION AX455554  
 VERSION AX455554.1 GI:21714622  
 KEYWORDS synthetic construct.  
 SOURCE synthetic construct.  
 ORGANISM artificial sequences.  
 REFERENCE 1  
 AUTHORS Bauer, S., Lipford, G. and Wagner, H.  
 TITLE Process for high throughput screening of cpg-based immuno-agonist/antagonist  
 JOURNAL Patent: WO 0222809-A 31 21-MAR-2002;  
 Coley Pharmaceutical GmbH (DE)

## BASE COUNT

BASE COUNT 2 a 4 c 14 g 0 t  
 ORIGIN

QY 1 AGCGGGGCGAGCGGGGCG 20  
 |||||  
 Db 1 AGCGGGGCGAGCGGGGCG 20

## RESULT 5

LOCUS AX472520 30 bp DNA linear PAT 09-AUG-2002  
 DEFINITION Sequence 15 from Patent WO02052039.  
 ACCESSION AX472520  
 VERSION AX472520.1 GI:22207424  
 KEYWORDS synthetic construct.  
 SOURCE synthetic construct.  
 ORGANISM artificial sequences.  
 REFERENCE 1  
 AUTHORS Blais, Y., Rousseau, P., Leblanc, B. and Camato, R.N.  
 TITLE Methods for selecting and producing selective pharmaceutical compounds and compositions using an established genetically altered cell-based library responsive to transcription factors; genetic constructs and library therefor  
 JOURNAL Patent: WO 02052039-A 15 04-JUL-2002;  
 Geneka Biotechnology Inc. (CA)

FEATURES  
Location/Qualifiers

LOCUS AX472520 30 bp DNA linear PAT 09-AUG-2002  
 DEFINITION Sequence 15 from Patent WO02052039.  
 ACCESSION AX472520  
 VERSION AX472520.1 GI:22207424  
 KEYWORDS synthetic construct.  
 SOURCE synthetic construct.  
 ORGANISM artificial sequences.  
 REFERENCE 1  
 AUTHORS Blais, Y., Rousseau, P., Leblanc, B. and Camato, R.N.  
 TITLE Methods for selecting and producing selective pharmaceutical compounds and compositions using an established genetically altered cell-based library responsive to transcription factors; genetic constructs and library therefor  
 JOURNAL Patent: WO 02052039-A 15 04-JUL-2002;  
 Geneka Biotechnology Inc. (CA)

## BASE COUNT

BASE COUNT 5 a 7 c 17 g 1 t  
 ORIGIN

QY 1 AGCGGGGCGAGCGGGGCG 20  
 |||||  
 Db 1 AGCGGGGCGAGCGGGGCG 20

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Db 7 AGCGGGGGCGAGCGGGGCG 26
|||||
AX476834 30 bp DNA linear PAT 12-AUG-2002
LOCUS
DEFINITION Sequence 11 from Patent WO02052037.
ACCESSION AX476834
VERSION AX476834.1 GI:22216110
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE
1 Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.
AUTHORS Method for screening and/or identifying factors that bind to
TITLE nucleic acids
JOURNAL Patent: WO 02052037-A 11 04-JUL-2002;
Geneka Biotechnology Inc. (CA)
FEATURES
1 Location/Qualifiers
source
1..30
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="NABE-probes"
BASE COUNT 5 a 7 c 17 g 1 t
ORIGIN
Query Match 100.0%; Score 20; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGCGGGGGCGAGCGGGGCG 20
|||||
Db 7 AGCGGGGGCGAGCGGGGCG 26
|||||
AX476857 30 bp DNA linear PAT 12-AUG-2002
LOCUS
DEFINITION Sequence 34 from Patent WO02052037.
ACCESSION AX476857
VERSION AX476857.1 GI:22216133
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE
1 Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.
AUTHORS Method for screening and/or identifying factors that bind to
TITLE nucleic acids
JOURNAL Patent: WO 02052037-A 34 04-JUL-2002;
Geneka Biotechnology Inc. (CA)
FEATURES
1 Location/Qualifiers
source
1..30
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="Double stranded NABE"
BASE COUNT 5 a 7 c 17 g 1 t
ORIGIN
Query Match 100.0%; Score 20; DB 6; Length 30;
Best Local Similarity 100.0%; Pred. No. 2e+03;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 AGCGGGGGCGAGCGGGGCG 20
|||||
Db 7 AGCGGGGGCGAGCGGGGCG 26
|||||
AX128389 47 bp DNA linear PAT 15-MAY-2001
LOCUS
DEFINITION Sequence 50 from Patent WO0130843.
ACCESSION AX128389
VERSION AX128389.1 GI:14134897
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE
1 (bases 1 to 47)
AUTHORS Barbas,C.F., Kadan,M. and Beerli,R.
TITLE Ligand activated transcriptional regulator proteins
JOURNAL Patent: WO 0130843-A 50 03-MAY-2001;
Novartis AG (CH); The Scripps Research Institute (US)
FEATURES
1 Location/Qualifiers
source
1..47
/organism="synthetic construct"
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JOURNAL Patent: WO 0140521-A 1611 07-JUN-2001;
FEATURES Curagen Corporation (US)
source Location/Qualifiers
1..51
/organism="Homo sapiens"
/db_xref="taxon:9606"
misc_feature 26
/note="1 of 2 allelic variants (1612 is other entry)
Accession number cg32120097"
BASE COUNT 6 a 22 c 15 g 8 t
ORIGIN

Query Match 71.0%; Score 14.2; DB 6; Length 51;
Best Local Similarity 84.2%; Pred. No. 2.2e+05;
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 AGCGGGGGCGAGCGGGGC 19
| | | | | | | | | | | | | | | |
Db 43 AGCGGGGGCCAGCGGGAGC 25

Search completed: December 12, 2002, 02:55:54
Job time : 323.116 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2002 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:07:11 ; Search time 815.797 Seconds  
(without alignments)  
397.047 Million cell updates/sec

Title: US-09-355-254F-8

Perfect score: 20

Sequence: 1 gattgcctgacgtcagag 20

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched: 16154066 seqs, 8097743376 residues

Total number of hits satisfying chosen parameters: 357874

Minimum DB seq length: 0

Maximum DB seq length: 100

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:\*\*

1: em\_estba:\*\*

2: em\_esthum:\*\*

3: em\_estlin:\*\*

4: em\_estmu:\*\*

5: em\_estov:\*\*

6: em\_estpl:\*\*

7: em\_estro:\*\*

8: em\_htc:\*\*

9: gb\_estl:\*\*

10: gb\_est2:\*\*

11: gb\_htc:\*\*

12: gb\_est3:\*\*

13: gb\_est4:\*\*

14: gb\_est5:\*\*

15: em\_estfun:\*\*

16: em\_estom:\*\*

17: gb\_gss:\*\*

18: em\_gss\_hum:\*\*

19: em\_gss\_inv:\*\*

20: em\_gss\_pln:\*\*

21: em\_gss\_vrt:\*\*

22: em\_gss\_fun:\*\*

23: em\_gss\_mam:\*\*

24: em\_gss\_mus:\*\*

25: em\_gss\_other:\*\*

26: em\_gss\_pro:\*\*

27: em\_gss\_rod:\*\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
C 1	14.2	71.0	78	17	AZ871578
C 2	13.4	67.0	50	17	AZ492887
C 3	13.2	66.0	49	17	BH861707
C 4	13.2	66.0	50	9	AU107195
C 5	13.2	66.0	67	9	AA721395
C 6	13.2	66.0	71	9	AA702828

C 7	13.2	66.0	91	17	AZ512653
C 8	13.2	66.0	93	9	AA560086
C 9	13.2	66.0	99	12	BF458655
C 10	12.8	64.0	34	17	AQ073796
C 11	12.8	64.0	40	9	AA848120
C 12	12.8	64.0	50	9	AU105994
C 13	12.8	64.0	69	17	BH256492
C 14	12.8	64.0	86	9	AI204909
C 15	12.8	64.0	88	9	AU260302
C 16	12.8	64.0	99	14	F26219
C 17	12.6	63.0	73	10	BE095402
C 18	12.6	63.0	73	17	BH862484
C 19	12.6	63.0	81	9	AA580289
C 20	12.6	63.0	84	10	AV952690
C 21	12.6	63.0	84	17	AZ312968
C 22	12.6	63.0	91	17	AZ494456
C 23	12.6	63.0	95	17	AZ810823
C 24	12.6	63.0	98	17	BH853467
C 25	12.4	62.0	93	14	W31755
C 26	12.4	62.0	94	17	AZ763191
C 27	12.2	61.0	43	17	BH796799
C 28	12.2	61.0	50	9	AU107190
C 29	12.2	61.0	50	9	AU107194
C 30	12.2	61.0	50	9	AU107198
C 31	12.2	61.0	54	17	AZ776635
C 32	12.2	61.0	55	17	AZ920728
C 33	12.2	61.0	64	9	AI253600
C 34	12.2	61.0	64	13	BI698792
C 35	12.2	61.0	74	9	AA218675
C 36	12.2	61.0	79	9	AA600457
C 37	12.2	61.0	80	9	AA985780
C 38	12.2	61.0	82	14	W05256
C 39	12.2	61.0	83	9	AA075933
C 40	12.2	61.0	85	9	AA716566
C 41	12.2	61.0	88	17	AZ430802
C 42	12.2	61.0	90	12	BG400826
C 43	12.2	61.0	97	14	H28506
C 44	12.2	61.0	100	13	BI002964
C 45	12.2	61.0	100	17	AZ780127

ALIGNMENTS

RESULT 1

AZ871578/C

LOCUS

DEFINITION

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

AZ871578, 78 bp DNA linear GSS 21-FEB-2001  
2M0184G04R Mouse 10kb plasmid UUGC1M library Mus musculus genomic  
clone UUGC2M0184G04 R, DNA sequence.  
AZ871578 GI:13077918  
GSS.  
house mouse.  
Mus musculus  
Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Mus.  
Dunn,D., Aoyagi,A., Barber,M., Beacorn,T., Duval,B., Hamil,C.,  
Islam,H., Longacre,S., Mahmoud,M., Meenen,E., Pedersen,T., Reilly  
,M., Rose,M., Rose,R., Stokes,R., Tingey,A., von Niederhausern,A.  
and Wright,D.,Weiss,R.  
Mouse whole genome scaffolding with paired end reads from 10kb  
plasmid inserts  
Unpublished (2000)  
Contact: Robert B. Weiss  
University of Utah Genome Center  
University of Utah  
Rm. 308, Biomedical Polymers Research Bldg., 20 S. 2030 E., SLIC, UT  
84112, USA  
Tel: 801 585 5606  
Fax: 801 585 7177  
Email: ddunn@genetics.utah.edu  
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FEATURES
  source
    Class: TDNA tagged.
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        /organism="Arabidopsis thaliana"
        /strain="Columbia 0"
        /db_xref="taxon:3702"
        /clone="SALK_087868"
        /note="PCR was performed on Arabidopsis thaliana lines
        each of which contains one or more TDNA insertion
        elements. The resultant fragment for each line was
        directly sequenced to determine the genomic sequence at
        the site of insertion. Details of the protocols used can
        be found at: http://signal.salk.edu/tdna_protocols.html"
BASE COUNT      16 a   14 c   7 g   12 t
ORIGIN
Query Match      66.0%; Score 13.2; DB 17; Length 49;
Best Local Similarity 83.3%; Pred. No. 2.8e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGCTCAGAGA 19
    ||| | ||||| |||||
Db 36 ATTCTTGACGTCAGAGA 19

RESULT 4
LOCUS      AU107195/c
DEFINITION AU107195 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone
ACCESSION  HEP21410, mRNA sequence.
VERSION     AU107195
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
REFERENCE    1 (bases 1 to 50)
AUTHORS     Suzuki.Y., Taira.H., Tsunoda.T., Mizushima-Sugano.J., Sese,J., Hata
            Y., Nakamura.Y., Suyama,A. and Sugano.S.
TITLE       Diverse transcriptional initiation revealed by fine, large-scale
            mapping of mRNA start sites
JOURNAL     EMBO Rep. 2 (5), 388-393 (2001)
MEDLINE     21270072
COMMENT     Contact: Yutaka Suzuki
            Department of Virology
            Institute of Medical Science, University of Tokyo
            4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan
            Email: ysuzuki@ims.u-tokyo.ac.jp
            Suzuki.Y., Yoshitomo-Nakagawa.K., Maruyama.K., Suyama.A. and Sugano
            S. Construction and characterization of a full length-enriched and
            a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

FEATURES
  source
    Location/Qualifiers
      1..50
        /organism="Homo sapiens"
        /db_xref="taxon:9606"
        /clone="HEP21410"
        /note="Differential display comparison of untreated and
        dimethylfumarate treated U937 cells"
BASE COUNT      12 a   14 c   14 g   10 t
ORIGIN
Query Match      66.0%; Score 13.2; DB 9; Length 50;
Best Local Similarity 83.3%; Pred. No. 2.8e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TTCCCTGACGTCAGAG 20
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Db 41 TTGGCTGACGTCACAG 24

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```

RESULT 5
LOCUS      AA721395/c
DEFINITION n273g08.s1 NCI_CGAP_GCB1 Homo sapiens cDNA clone IMAGE:1301150 3',
            mRNA sequence.
ACCESSION  AA721395
VERSION     AA721395.1
KEYWORDS    EST.
SOURCE      human.
ORGANISM    Homo sapiens
REFERENCE    1 (bases 1 to 67)
AUTHORS     NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
            National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
            Tumor Gene Index
            Unpublished (1997)
            Contact: Robert Strausberg, Ph.D.
            Email: cgapbs-r@mail.nih.gov
            Tissue Procurement: Louis M. Staudt, M.D., Ph.D., David Allman,
            Ph.D., Gerald Marti, M.D.
            CDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima
            Bonaldo, Ph.D.
            CDNA Library Arrayed by: Greg Lennon, Ph.D.
            DNA Sequencing by: Washington University Genome Sequencing Center
            Clone distribution: NCI-CGAP clone distribution information can be
            found through the I.M.A.G.E. Consortium/LLNL at:
            www.bio.llnl.gov/bbrp/image/image.html
            Insert length: 835 Std Error: 0.00
            Seq primer: -40ml3 fwd. ET from Amersham
            High quality sequence stop: 57.
FEATURES
  Location/Qualifiers
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      /db_xref="taxon:9606"
      /clone="IMAGE:1301150"
      /clone_lib="NCI_CGAP_GCB1"
      /tissue_type="germinal center B cell"
      /lab_host="DH10B"
      /note="Vector: pT73D-Pac (Pharmacia) with a modified
      polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
      was prepared from human tonsillar cells enriched for
      germinal center B cells by flow sorting (CD20+, IgD-),
      provided by Dr. Louis M. Staudt (NCI), Dr. David Allman
      (NCI) and Dr. Gerald Marti (CBER). CDNA synthesis was
      primed with a Not I - oligo(dT) primer
      15'-TGTTACCAATCTGAAGTGGAGCGGCCCTCATTTTTTTTTTTTTTTT-3'
      1. Double-stranded cDNA was ligated to Eco RI adaptors
      (Pharmacia), digested with Not I and cloned into the Not I
      and Eco RI sites of the modified pT73 vector. Library
      went through one round of normalization, and was
      constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT      14 a   24 c   15 g   14 t
ORIGIN
Query Match      66.0%; Score 13.2; DB 9; Length 67;
Best Local Similarity 83.3%; Pred. No. 3.2e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 18
    ||||| |||| |
Db 30 GATTGCTGACCCAGAG 13

RESULT 6
LOCUS      AA702828
DEFINITION z177b06.s1 Soares fetal_liver_spleen_LNFLS_S1 Homo sapiens cDNA
            clone IMAGE:436787 3' similar to contains element CER repetitive
            element ;, mRNA sequence.
ACCESSION  AA702828
VERSION     AA702828.1

```



# TITLE JOURNAL COMMENT

The WashU-HMI Mouse EST Project  
Unpublished (1996)  
Contact: Marra M/Mouse EST Project  
WashU-HMI Mouse EST Project  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810  
Email: mouseest@wustl.edu  
This clone is available royalty-free through LILNL; contact the  
IMAGE Consortium (info@image.lilnl.gov) for further information.  
MGI:554368  
Seq primer: -28ml3 rev1 ET from Amersham  
High quality sequence stop: 85.

## FEATURES source

1. .93  
Location/Qualifiers  
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/db\_xref="taxon:10090"  
/clone="IMAGE:973640"  
/clone\_lib="Stratagene mouse Tcell 937311"  
/tissue\_type="Tcell"  
/dev\_stage="M30 CD4+ cells"  
/lab\_host="SOLR (kanamycin resistant)"  
/note="Organ: blood; Vector: pBluescript SK-; Site\_1:  
EcoRI; Site\_2: XhoI; Cloned unidirectionally. Primer:  
Oligo dT. M30 CD4+ cells. Average insert size: 1.0 Kb;  
Uni-ZAP XR Vector; -5' adaptor sequence: 5' GAATTCGCGCAGG  
3' -3' adaptor sequence: 5' CTCGAGTGTGTTTTTTTTTTT 3'

BASE COUNT 22 a 17 c 28 g 26 t  
ORIGIN  
Query Match 66.0%; Score 13.2; DB 9; Length 93;  
Best Local Similarity 83.3%; Pred. No. 3.6e+04;  
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 2 ATTCCTGACGTCAGAGA 19  
Db 62 ATGCGTGTGATGTCAGAGA 79  
||||| ||| |||||

## RESULT 9 BF458655/c

LOCUS BF458655 99 bp mRNA linear EST 01-DEC-2000  
DEFINITION UI-M-B21-b1t-g-01-0-UI.s1 NIH\_BMAP\_MHI2\_S1 Mus musculus cDNA clone  
UI-M-B21-b1t-g-01-0-UI 3', mRNA sequence.  
ACCESSION BF458655  
VERSION BF458655.1 GI:11524824  
KEYWORDS EST.  
SOURCE house mouse.  
ORGANISM Mus musculus  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
REFERENCE 1 (bases 1 to 99)  
Bonaldo,M.F., Lennon,G. and Soares,M.B.  
Discovery Normalization and subtraction: two approaches to facilitate gene  
Genome Res. 6 (9), 791-806 (1996)  
9704477  
COMMENT Contact: Chin, H  
National Institute of Mental Health  
6001 Executive Blvd. Room 7N-7190, MSC 9643, Bethesda, MD  
20892-9643, USA  
Tel: 301 443 1706  
Fax: 301 443 9890  
Email: mEST@mail.nih.gov

The sequence contained an oligo-dT track that was present in the  
oligonucleotide that was used to prime the synthesis of first  
strand cDNA and therefore this may represent a bonafide poly A  
tail. The sequence tag present in the cDNA between the NotI site  
and the oligo-dT track served to verify it as a clone from the  
hippocampus tissue cDNA Library Preparation: M.B. Soares Lab Clone  
distribution: Researchers may obtain BMAP cDNA clones from RESEARCH  
GENETICS. It should be noted that Bento Soares is generating a

small number of additional specialized non-redundant arrays of BMAP  
cDNAs whose availability will be considered under appropriate and  
limited collaborative arrangements The following repetitive  
elements were found in this cDNA sequence: 1-21,  
>AT-rich#Low complexity  
Seq primer: M13 Forward  
POLYA=yes.

## FEATURES source

Location/Qualifiers  
1. .99  
/organism="Mus musculus"  
/strain="C57BL/6J"  
/db\_xref="taxon:10090"  
/clone="UI-M-B21-b1t-g-01-0-UI"  
/clone\_lib="NIH\_BMAP\_MHI2\_S1"  
/dev\_stage="27-32 days"  
/lab\_host="DH10B (Life Technologies)"  
/note="Vector: pT73D-Pac (Pharmacia) with a modified  
polylinker; Site\_1: Not I; Site\_2: Eco RI; The  
NIH\_BMAP\_MHI2\_S1 library is a subtracted library derived  
from NIH\_BMAP\_MHI2. NIH\_BMAP\_MHI2 is a library derived  
from mouse hippocampus tissue. For a detailed description  
of the library from which this clone was derived, please  
visit our web site at brainest.eng.uiowa.edu.  
TAG\_LIB=NIH\_BMAP\_MHI2\_S1  
TAG\_TISSUE=hippocampus  
TAG\_SEQ=TAGCC"

BASE COUNT 18 a 26 c 32 g 23 t  
ORIGIN

Query Match 66.0%; Score 13.2; DB 12; Length 99;  
Best Local Similarity 83.3%; Pred. No. 3.7e+04;  
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

OY 3 TTGCCTGACGTCAGAGAG 20  
Db 57 TTCCCTGGGTGAGAGAG 40  
||||| ||| |||||

## RESULT 10 AQ073796/c

LOCUS AQ073796 34 bp DNA linear GSS 23-AUG-2000  
DEFINITION EP(3)379 Drosophila melanogaster Ep line Drosophila melanogaster  
genomic sequence recovered from Both 5' and 3' ends of P element,  
DNA sequence.  
ACCESSION AQ073796  
VERSION AQ073796.1 GI:3403838  
KEYWORDS GSS.  
SOURCE fruit fly.  
ORGANISM Drosophila melanogaster  
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota;  
Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha;  
Ephydroidea; Drosophilidae; Drosophila.  
REFERENCE 1 (bases 1 to 34)  
Liao,G.-C., Rehm,E.J. and Rubin,G.M.  
Insertion site preferences of the P transposable element in  
Drosophila melanogaster  
Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3347-3351 (2000)  
20202638  
COMMENT Contact: Gerald Rubin  
Berkeley Drosophila Genome Project  
University of California, Berkeley  
LSA Building, Berkeley, CA 94720-3200, USA  
Fax: 5106439947  
Email: gerry@fruitfly.berkeley.edu  
Sequence recovery method was inverse PCR.  
Sequence orientation is forward strand relative to 5' end of P  
element

The P element insertion position is base 15 in the 34 bases. This  
insertion position refers to the first base of the 8 base target  
recognition sequence.  
Class: transposon-tagged.

FEATURES  
source

Location/Qualifiers  
1. .34  
/organism="Drosophila melanogaster"  
/db\_xref="taxon:7227"  
/clone\_lib="Drosophila melanogaster EP line"  
/note="Inverse PCR was performed on Drosophila melanogaster strains each of which contains a single EP transposable element insertion. (The generation of these insertion strains is described in Rorth P, Szabo K, Bailey A, Lavery T, Rehm J, Rubin GM, Weigmann K, Milan M, Benes V, Ansorge W, Cohen SM. 1998. Systematic gain-of-function genetics in Drosophila. Development 6:1049-1057.) The resultant fragment for each strain was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at [http://fruitfly.berkeley.edu/p\\_disrupt/inverse\\_pcr.html](http://fruitfly.berkeley.edu/p_disrupt/inverse_pcr.html)."

4 a 11 c 10 g 9 t

BASE COUNT  
ORIGIN

Query Match 64.0%; Score 12.8; DB 17; Length 34;  
Best Local Similarity 87.5%; Pred. No. 3.6e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 GCGTGACGTGAGAG 20

DB 32 GCGTGACGTGAGAG 17

RESULT 11  
AA848120/c

LOCUS  
DEFINITION  
oe05a03.s1 NCI\_CGAP\_Ov2 Homo sapiens cDNA clone IMAGE:1384972 similar to gb:M35663 INTERFERON-INDUCED, DOUBLE-STRANDED RNA-ACTIVATED PROTEIN KINASE (HUMAN);, mRNA sequence.

ACCESSION  
AA848120

VERSION  
EST.

KEYWORDS  
human.

SOURCE  
Homo sapiens

ORGANISM  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE  
Mammalia; Euthera; Primates; Catarrhini; Homiidae; Homo.

1 (bases 1 to 40)

NCI-CGAP <http://www.ncbi.nlm.nih.gov/ncicgap>.

National Cancer Institute, Cancer Genome Anatomy Project (CGAP),

Tumor Gene Index

Unpublished (1997)

Contact: Robert Strausberg, Ph.D.

Email: [cgaps@mail.nih.gov](mailto:cgaps@mail.nih.gov)

Tissue Procurement: Christopher A. Moskaluk, M.D., Michael R.

Emmert-Buck, M.D., Ph.D.

cDNA Library Preparation: David B. Krizman, Ph.D.

DNA Sequencing by: Washington University Genome Sequencing Center

Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at:

[www-bio.llnl.gov/bbrp/image/image.html](http://www-bio.llnl.gov/bbrp/image/image.html)

Trace considered overall poor quality

Insert Length: 1464 Std Error: 0.00

Seq primer: -40m13 fwd. ET from Amersham

High quality sequence stop: 1.

Location/Qualifiers

1. .40

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="IMAGE:1384972"

/clone\_lib="NCI\_CGAP\_Ov2"

/sex="female"

/tissue\_type="ovary"

/lab\_host="DH10B"

/note="Vector: pAMP10; mRNA made from invasive ovarian tumor, cDNA made by oligo-dT priming. Non-directionally cloned. Size-selected on agarose gel, average insert size

600 bp. Reference: Krizman et al. (1996) Cancer Research 56:5380-5383."

8 a 14 c 9 g 9 t

BASE COUNT  
ORIGIN

Query Match 64.0%; Score 12.8; DB 9; Length 40;  
Best Local Similarity 87.5%; Pred. No. 3.9e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTGAG 16

DB 40 GATTGCTGACGTGAG 25

RESULT 12  
AU105994/c

LOCUS  
DEFINITION  
AU105994 Sugano Homo sapiens cDNA library Homo sapiens cDNA clone KATA2602, mRNA sequence.

ACCESSION  
AU105994

VERSION  
AU105994.1

KEYWORDS  
EST.

SOURCE  
human.

ORGANISM  
Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

REFERENCE  
Mammalia; Euthera; Primates; Catarrhini; Homiidae; Homo.

1 (bases 1 to 50)

AUTHORS  
Suzuki.Y., Taira.H., Tsunoda.T., Mizushima-Sugano.J., Sese.J., Hata

, Y., Ota.T., Isogai.T., Tanaka.T., Morishita.S., Okubo.K., Sakaki

, Y., Nakamura.Y., Suyama.A. and Sugano.S.

Diverse transcriptional initiation revealed by fine, large-scale

mapping of mRNA start sites

EMBO Rep. 2 (5), 388-393 (2001)

21270072

CONTACT: Yutaka Suzuki

Department of Virology

Institute of Medical Science, University of Tokyo

4-6-1, Shirokanedai, Minatoku, Tokyo 108-8639, Japan

Email: [ysuzuki@ims.u-tokyo.ac.jp](mailto:ysuzuki@ims.u-tokyo.ac.jp)

Suzuki.Y., Yoshitomo-Nakagawa.K., Maruyama.K., Suyama.A. and Sugano

, S. Construction and characterization of a full length-enriched and

a 5'-end-enriched cDNA library. Gene 200 (1-2), 149-156 (1997).

Location/Qualifiers

1. .50

/organism="Homo sapiens"

/db\_xref="taxon:9606"

/clone="KATA2602"

/clone\_lib="Sugano Homo sapiens cDNA library"

/note="Differential display comparison of untreated and

dimethylfumarate treated U937 cells"

BASE COUNT 11 a 19 c 9 g 11 t

BASE COUNT  
ORIGIN

Query Match 64.0%; Score 12.8; DB 9; Length 50;  
Best Local Similarity 87.5%; Pred. No. 4.3e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTGAG 16

DB 29 GATTGCTGACGTGAG 14

RESULT 13  
BH256492

LOCUS  
DEFINITION  
BH256492  
KG03686-5prime Drosophila melanogaster PISUPor-P] p element insertion lines Drosophila melanogaster genomic Sequence recovered from 5' end of P element, DNA sequence.

ACCESSION  
BH256492

VERSION  
BH256492.1

KEYWORDS  
GSS.

SOURCE  
fruit fly.

ORGANISM  
Drosophila melanogaster

Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.

REFERENCE  
AUTHORS  
1 (bases 1 to 69)  
Levis, R., Hoskins, R., Liao, G., Mozen, N., Tsang, G., He, Y., Karpen, G., Bellen, H., Rubin, G. and Spradling, A.  
The Berkeley Drosophila Genome Project Gene Disruption Project  
Unpublished (2001)  
CONTACT: Gerald Rubin  
Berkeley Drosophila Genome Project  
University of California, Berkeley  
LSA Building, Berkeley, CA 94720-3200, USA  
Fax: 5106433947

EMAIL: gerry@fruitfly.berkeley.edu  
Sequence recovery method was inverse PCR.  
Sequence orientation is forward strand relative to 5' end of P element

The P element insertion position is base 62 in the 69 bases. This insertion position refers to the first base of the 8 base target recognition sequence.  
Class: transposon-tagged.  
Location/Qualifiers

FEATURES  
source  
1..69  
/organism="Drosophila melanogaster"  
/db\_xref="taxon:7227"  
/clone\_lib="Drosophila melanogaster P{SUPor-P} P element insertion lines"

/note="Inverse PCR was performed on Drosophila melanogaster strains each of which contains one or more P{SUPor-P} P-element transposon insertion. The resultant fragment for each strain was directly sequenced to determine the genomic sequence at the site of insertion. Details of the protocols used can be found at <http://www.fruitfly.org/about/methods/inverse.pcr.html>."

BASE COUNT 15 a 15 c 18 g 21 t

ORIGIN  
Query Match 64.0%; Score 12.8; DB 17; Length 69;  
Best Local Similarity 87.5%; Pred. No. 4.9e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 ATTGCCTGAGTCGACA 17  
||||| III |||||  
Db 2 ATTCGCTGAAGTCGACA 17

RESULT 14  
AI204909  
LOCUS  
AI204909 86 bp mRNA linear EST 15-OCT-1998  
DEFINITION  
an05g11.x1 Stratagene schizo brain S11 Homo sapiens cDNA clone  
IMAGE:1684772 3' similar to gb:M55053 CYTOCHROME P450 1A2 (HUMAN);,  
mRNA sequence.

ACCESSION  
AI204909  
VERSION  
AI204909.1 GI:3757971  
KEYWORDS  
EST.  
SOURCE  
Homo sapiens

ORGANISM  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
1 (bases 1 to 86)  
Miller, L., Allen, M., Bowles, L., Dubuque, T., Geisel, G., Jost, S.,  
Krizman, D., Kucaba, T., Lacy, M., Le, N., Lennon, G., Marra, M., Martin,  
J., Moore, B., Schellenberg, K., Steptoe, M., Tan, F., Theising, B.,  
White, Y., Wylie, T., Waterston, R. and Wilson, R.

WashU-NCI human EST Project  
Unpublished (1997)  
Contact: Wilson RK  
Washington University School of Medicine  
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108  
Tel: 314 286 1800  
Fax: 314 286 1810

EMAIL: est@watson.wustl.edu  
This clone is available royalty-free through LLNL ; contact the

IMAGE Consortium ([info@image.llnl.gov](mailto:info@image.llnl.gov)) for further information.  
Trace considered overall poor quality  
Seq primer: -40UP from Gibco  
High quality sequence stop: 1.

FEATURES  
source  
1..86  
Location/Qualifiers  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/clone="IMAGE:1684772"  
/clone\_lib="Stratagene schizo brain S11"  
/sex="male"  
/tissue\_type="schizophrenic brain S-11 frontal lobe"  
/dev\_stage="34 years old"  
/lab\_host="SOLR (kanamycin resistant)"  
/note="Vector: Bluescript SK-; Site:1: EcoRI; Library constructed from S-11 frontal lobe, male, 34 years old, 50% caucasian, 50% Aleutian. Schizophrenic suicide. Random primed into EcoRI site of ZAP II Vector. Mass excised. Custom library. Avg insert length 1.4kb. Material obtained by Johnston N., Torrey, E.F., Yolken R., and the Stanley Neuropathology Consortium - Analysis of RNAs from the Brains of Individuals with Psychiatric Diseases (Unpublished) Stanley Neurovirology Laboratory, Johns Hopkins School of Medicine, Baltimore MD."

BASE COUNT 26 a 19 c 24 g 17 t

ORIGIN  
Query Match 64.0%; Score 12.8; DB 9; Length 86;  
Best Local Similarity 87.5%; Pred. No. 5.4e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 GCCTGACGTCAGAGAG 20  
||||| III |||||  
Db 43 GCCTGCGCAGAGAG 58

RESULT 15  
AU260302/c  
LOCUS  
AU260302 88 bp mRNA linear EST 25-APR-2002  
DEFINITION  
AU260302 3'-directed mouse cDNA library Mus musculus cDNA clone  
BED0016644 3', mRNA sequence.

ACCESSION  
AU260302  
VERSION  
AU260302.1 GI:20327661  
KEYWORDS  
EST.  
SOURCE  
house mouse.  
Mus musculus

ORGANISM  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.  
1 (bases 1 to 88)  
Kato, K. and Matoba, R.

REFERENCE  
AUTHORS  
Kato, K. and Matoba, R.  
TITLE  
Generation of expressed sequence tags from mouse brain  
JOURNAL  
Unpublished (2002)  
COMMENT  
Contact: Kikuya Kato  
Graduate School of Biological Sciences  
Nara Institute of Science and Technology  
8916-5 Takayama, Ikoma, Nara 630-0101, Japan  
Tel: 81-743-72-5581  
Fax: 81-743-72-5589  
Email: [kkato@bs.nara.ac.jp](mailto:kkato@bs.nara.ac.jp),  
URL: <http://love2.aist-nara.ac.jp/BED/index.html>.

LOCATION/Qualifiers  
1..88  
/organism="Mus musculus"  
/db\_xref="taxon:10090"  
/clone="BED0016644"

/clone\_lib="3'-directed mouse cDNA library"  
/tissue\_type="brain"  
/note="Vector: pGEM-T-easy"  
BASE COUNT 27 a 26 c 19 g 16 t

ORIGIN  
Query Match 64.0%; Score 12.8; DB 9; Length 88;  
Best Local Similarity 87.5%; Pred. No. 5.4e+04;

---

Matches	14;	Conservative	0;	Mismatches	2;	Indels	0;	Gaps	0;
Qy	5	GCCTGACGTCAGAGAG	20						
Db	19	GCCTGAACCTCAGAGAG	4						

Search completed: December 12, 2002, 06:03:49  
Job time : 834.154 secs

Result NO.	Score	Query Match	Length	DB	ID	Description
1	20	100.0	27	10	US-09-878-918-2	Sequence 2, Appli
2	19	95.0	23	10	US-09-816-763-148	Sequence 148, App
3	12.8	64.0	50	10	US-09-875-945-4	Sequence 4, Appli
4	12.8	64.0	97	10	US-09-864-761-19704	Sequence 19704, A
5	12.6	63.0	57	10	US-09-969-193-60	Sequence 60, Appl
c	12.6	63.0	57	10	US-09-969-193-61	Sequence 61, Appl
7	12.6	63.0	63	10	US-09-969-192-58	Sequence 58, Appl
c	12.6	63.0	86	10	US-09-983-503-5019	Sequence 5019, Ap
9	12.2	61.0	24	10	US-09-420-433-62	Sequence 62, Appl
c	12.2	61.0	24	10	US-09-784-911-30	Sequence 30, Appl
11	12.2	61.0	99	10	US-09-769-066-11	Sequence 11, Appl
12	12	60.0	63	10	US-09-998-598-2339	Sequence 2339, Ap
13	12	60.0	86	9	US-10-040-497-79	Sequence 79, Appl
14	12	60.0	92	10	US-09-864-761-28968	Sequence 28968, A
15	11.8	59.0	18	8	US-08-983-605-140	Sequence 140, App
16	11.6	58.0	18	10	US-09-303-510-50	Sequence 50, Appl
17	11.6	58.0	18	10	US-09-303-040-50	Sequence 50, Appl
c	11.6	58.0	20	9	US-10-060-301-123	Sequence 123, App
c	11.6	58.0	41	10	US-09-238-351-29	Sequence 29, Appl

; TITLE OF INVENTION: DETECTION AND/OR THE QUANTIFICATION OF TRANSCRIPTIONAL

; FILE REFERENCE: VANM212.001AUS  
; CURRENT APPLICATION NUMBER: US/09/816,763  
; PRIOR FILING DATE: 2001-03-23  
; PRIOR APPLICATION NUMBER: EP 00870057.7  
; PRIOR FILING DATE: 2000-03-24  
; NUMBER OF SEQ ID NOS: 150  
; SOFTWARE: FastSeq for Windows version 4.0  
; SEQ ID NO 148  
; LENGTH: 23  
; TYPE: DNA  
; ORGANISM: Artificial Sequence  
; FEATURE:  
; OTHER INFORMATION: CREB consensus sequence  
US-09-816-763-148

Query Match 95.0%; Score 19; DB 10; Length 23;  
Best Local Similarity 100.0%; Pred. No. 0.62;  
Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGTCAGAG 20  
|||||  
DB 1 ATTGCCTGACGTCAGAG 19

## RESULT 3

US-09-875-945-4  
; Sequence 4, Application US/09875945  
; Patent No. US20020098169A1  
; GENERAL INFORMATION:  
; APPLICANT: METCON MEDICIN AB  
; APPLICANT: SMITH, Ulf  
; TITLE OF INVENTION: No. US20020098169A1el sequences and their use  
; FILE REFERENCE: 45513MH  
; CURRENT APPLICATION NUMBER: US/09/875,945  
; CURRENT FILING DATE: 2001-06-08  
; PRIOR APPLICATION NUMBER: SE 0002189-9  
; PRIOR FILING DATE: 2000-06-09  
; PRIOR APPLICATION NUMBER: US 60/210,207  
; PRIOR FILING DATE: 2000-06-08  
; NUMBER OF SEQ ID NOS: 14  
; SOFTWARE: PatentIn Ver. 2.1  
; SEQ ID NO 4  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
US-09-875-945-4

Query Match 64.0%; Score 12.8; DB 10; Length 50;  
Best Local Similarity 87.5%; Pred. No. 9.4e+02;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAG 16  
|||||  
DB 6 GATTGCTGACGTCAG 21

## RESULT 4

US-09-864-761-19704  
; Sequence 19704, Application US/09864761  
; Patent No. US20020048763A1  
; GENERAL INFORMATION:  
; APPLICANT: Penn, Sharron G.  
; APPLICANT: Rank, David R.  
; APPLICANT: Hanzel, David K.  
; TITLE OF INVENTION: HUMAN GENOME-DERIVED SINGLE EXON NUCLEIC ACID PROBES USEFUL FOR  
; FILE REFERENCE: Aeonica-X-1  
; CURRENT APPLICATION NUMBER: US/09/864,761  
; CURRENT FILING DATE: 2001-05-23  
; PRIOR APPLICATION NUMBER: US 60/180,312

; PRIOR FILING DATE: 2000-02-04  
; PRIOR APPLICATION NUMBER: US 60/207,456  
; PRIOR FILING DATE: 2000-05-26  
; PRIOR APPLICATION NUMBER: US 09/632,366  
; PRIOR FILING DATE: 2000-08-03  
; PRIOR APPLICATION NUMBER: GB 24263.6  
; PRIOR FILING DATE: 2000-10-04  
; PRIOR APPLICATION NUMBER: US 60/236,359  
; PRIOR FILING DATE: 2000-09-27  
; PRIOR APPLICATION NUMBER: PCT/US01/006666  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006667  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006664  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006669  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006665  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006668  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006663  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006662  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006661  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: PCT/US01/006670  
; PRIOR FILING DATE: 2001-01-30  
; PRIOR APPLICATION NUMBER: US 60/234,687  
; PRIOR FILING DATE: 2000-09-21  
; PRIOR APPLICATION NUMBER: US 09/608,408  
; PRIOR FILING DATE: 2000-06-30  
; PRIOR APPLICATION NUMBER: US 09/774,203  
; PRIOR FILING DATE: 2001-01-29  
; NUMBER OF SEQ ID NOS: 49117  
; SOFTWARE: Annonmax Sequence Listing Engine vers. 1.1  
; SEQ ID NO 19704  
; LENGTH: 97  
; TYPE: DNA  
; ORGANISM: Homo sapiens  
; FEATURE:  
; OTHER INFORMATION: MAP TO AC009490.4  
; OTHER INFORMATION: EXPRESSED IN FETAL LIVER, SIGNAL = 1.6  
; OTHER INFORMATION: EXPRESSED IN LUNG, SIGNAL = 1.8  
; OTHER INFORMATION: EXPRESSED IN BONE MARROW, SIGNAL = 1.5  
; OTHER INFORMATION: EXPRESSED IN HEPA, SIGNAL = 1.7  
; OTHER INFORMATION: EXPRESSED IN HEART, SIGNAL = 1.5  
; OTHER INFORMATION: EXPRESSED IN PLACENTA, SIGNAL = 1.9  
; OTHER INFORMATION: EXPRESSED IN BT474, SIGNAL = 1.6  
; OTHER INFORMATION: EXPRESSED IN ADULT LIVER, SIGNAL = 2  
; OTHER INFORMATION: EXPRESSED IN BRAIN, SIGNAL = 1.7  
; OTHER INFORMATION: EXPRESSED IN HBL100, SIGNAL = 1.7  
; OTHER INFORMATION: NT HIT: X56160.1, EVALUATE 3.20e+00  
; OTHER INFORMATION: EST\_HUMAN HIT: AV648669.1, EVALUATE 1.90e-02  
US-09-864-761-19704

Query Match 64.0%; Score 12.8; DB 10; Length 97;  
Best Local Similarity 87.5%; Pred. No. 1e+03;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 GCCTGACGTCAGAG 20  
|||||  
DB 74 GCAGACATCAGAG 89

## RESULT 5

US-09-969-192-60  
; Sequence 60, Application US/09969192  
; Patent No. US20020151027A1  
; GENERAL INFORMATION:  
; APPLICANT: WICKHAM, THOMAS J.  
; APPLICANT: ROELVINK, PETRUS W.



;; KOVESDI, IMRE  
;; TITLE OF INVENTION: TARGETING ADENOVIRUS WITH USE OF  
;; CONSTRAINED PEPTIDE MOTIFS  
;;  
;; NUMBER OF SEQUENCES: 80  
;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: Leydig, Voit & Mayer, Ltd.  
;; STREET: Two Prudential Plaza - 49th Floor  
;; CITY: Chicago  
;; STATE: Illinois  
;; COUNTRY: USA  
;; ZIP: 60601  
;;  
;; COMPUTER READABLE FORM:  
;; MEDIUM TYPE: Floppy disk  
;; COMPUTER: IBM PC compatible  
;; OPERATING SYSTEM: PC-DOS/MS-DOS  
;; SOFTWARE: PatentIn Release #1.0, Version #1.30  
;;  
;; CURRENT APPLICATION NUMBER: US/09/969,192  
;; FILING DATE: 01-Oct-2001  
;;  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 9-455061  
;; FILING DATE: 06-DEC-1999  
;; APPLICATION NUMBER: US 9-130225  
;; FILING DATE: 06-AUG-1998  
;; APPLICATION NUMBER: US 8-701124  
;; FILING DATE: 21-AUG-1996  
;;  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Hefner, M. Daniel  
;; REGISTRATION NUMBER: 41,826  
;; REFERENCE/DOCKET NUMBER: 213564  
;;  
;; INFORMATION FOR SEQ ID NO: 60:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 57 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: single  
;; TOPOLOGY: linear  
;;  
;; MOLECULE TYPE: other nucleic acid  
;; SEQUENCE DESCRIPTION: SEQ ID NO: 60:  
US-09-969-192-60

Query Match 63.0%; Score 12.6; DB 10; Length 57;  
Best Local Similarity 78.9%; Pred. No. 1.2e+03;  
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ATTGCCTGAGTCGAGAG 20  
| | | | | | | | | |  
Db 8 AATTCTTGAGTCGAGAG 26

RESULT 6  
US-09-969-192-61/c  
; Sequence 61, Application US/09969192  
; Patent No. US20020151027A1  
; GENERAL INFORMATION:  
; APPLICANT: WICKHAM, THOMAS J.  
; ROELVINK, PETRUS W.  
; KOVESDI, IMRE  
; TITLE OF INVENTION: TARGETING ADENOVIRUS WITH USE OF  
; CONSTRAINED PEPTIDE MOTIFS  
;  
; NUMBER OF SEQUENCES: 80  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Leydig, Voit & Mayer, Ltd.  
; STREET: Two Prudential Plaza - 49th Floor  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60601  
;  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:

;; APPLICATION NUMBER: US/09/969,192  
;; FILING DATE: 01-Oct-2001  
;;  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 9-455061  
;; FILING DATE: 06-DEC-1999  
;; APPLICATION NUMBER: US 9-130225  
;; FILING DATE: 06-AUG-1998  
;; APPLICATION NUMBER: US 8-701124  
;; FILING DATE: 21-AUG-1996  
;;  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Hefner, M. Daniel  
;; REGISTRATION NUMBER: 41,826  
;; REFERENCE/DOCKET NUMBER: 213564  
;;  
;; INFORMATION FOR SEQ ID NO: 61:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 57 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: single  
;; TOPOLOGY: linear  
;;  
;; MOLECULE TYPE: other nucleic acid  
;; SEQUENCE DESCRIPTION: SEQ ID NO: 61:  
US-09-969-192-61

Query Match 63.0%; Score 12.6; DB 10; Length 57;  
Best Local Similarity 78.9%; Pred. No. 1.2e+03;  
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ATTGCCTGAGTCGAGAG 20  
| | | | | | | | | |  
Db 54 AATTCTTGAGTCGAGAG 36

RESULT 7  
US-09-969-192-58  
; Sequence 58, Application US/09969192  
; Patent No. US20020151027A1  
; GENERAL INFORMATION:  
; APPLICANT: WICKHAM, THOMAS J.  
; ROELVINK, PETRUS W.  
; KOVESDI, IMRE  
; TITLE OF INVENTION: TARGETING ADENOVIRUS WITH USE OF  
; CONSTRAINED PEPTIDE MOTIFS  
;  
; NUMBER OF SEQUENCES: 80  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: Leydig, Voit & Mayer, Ltd.  
; STREET: Two Prudential Plaza - 49th Floor  
; CITY: Chicago  
; STATE: Illinois  
; COUNTRY: USA  
; ZIP: 60601  
;  
; COMPUTER READABLE FORM:  
; MEDIUM TYPE: Floppy disk  
; COMPUTER: IBM PC compatible  
; OPERATING SYSTEM: PC-DOS/MS-DOS  
; SOFTWARE: PatentIn Release #1.0, Version #1.30  
; CURRENT APPLICATION DATA:  
; APPLICATION NUMBER: US/09/969,192  
; FILING DATE: 01-Oct-2001  
; PRIOR APPLICATION DATA:  
; APPLICATION NUMBER: US 9-455061  
; FILING DATE: 06-DEC-1999  
; APPLICATION NUMBER: US 9-130225  
; FILING DATE: 06-AUG-1998  
; APPLICATION NUMBER: US 8-701124  
; FILING DATE: 21-AUG-1996  
; ATTORNEY/AGENT INFORMATION:  
; NAME: Hefner, M. Daniel  
; REGISTRATION NUMBER: 41,826  
; REFERENCE/DOCKET NUMBER: 213564  
; INFORMATION FOR SEQ ID NO: 58:  
; SEQUENCE CHARACTERISTICS:  
; LENGTH: 63 base pairs  
; TYPE: nucleic acid

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;
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; SEQUENCE DESCRIPTION: SEQ ID NO: 58:
US-09-969-192-58
Query Match 63.0%; Score 12.6; DB 10; Length 63;
Best Local Similarity 78.9%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 2 ATTGCCTGACGTCAGAG 20
Db 9 AATTCTTGACGTCGGAG 27

RESULT 8
US-09-983-965-5019/c
; Sequence 5019, Application US/09983965
; Patent No. US20020137160A1
; GENERAL INFORMATION:
; APPLICANT: Warren, Wesley C.
; APPLICANT: Tao, Nengbing
; APPLICANT: Byatt, John C.
; APPLICANT: Mathialagan, Nagappan
; TITLE OF INVENTION: NUCLEIC ACID AND OTHER MOLECULES ASSOCIATED WITH LACTATION AND
; FILE REFERENCE: 37-21(10297)C
; CURRENT APPLICATION NUMBER: US/09/983,965
; CURRENT FILING DATE: 2001-10-26
; PRIOR APPLICATION NUMBER: US 09/465,231
; PRIOR FILING DATE: 1999-12-15
; PRIOR APPLICATION NUMBER: US 60/113,678
; PRIOR FILING DATE: 1998-12-17
; NUMBER OF SEQ ID NOS: 5912
; SEQ ID NO 5019
; LENGTH: 86
; TYPE: DNA
; ORGANISM: Bos taurus
; FEATURE:
; OTHER INFORMATION: Clone ID: 29-LIB34-022-Q1-E1-H1
US-09-983-965-5019

Query Match 63.0%; Score 12.6; DB 10; Length 86;
Best Local Similarity 78.9%; Pred. No. 1.2e+03;
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAGA 19
Db 61 GATGGCTGGGTCGAAGA 43

RESULT 9
US-09-420-433-62
; Sequence 62, Application US/09420433
; Patent No. US20020096480A1
; GENERAL INFORMATION:
; APPLICANT: Sidransky, David
; TITLE OF INVENTION: NUCLEIC ACID MUTATION DETECTION IN
; TITLE OF INVENTION: HISTOLOGIC TISSUE
; NUMBER OF SEQUENCES: 82
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Spensley Horn Jubas & Lubitz
; STREET: 1880 Century Park East, Suite 500
; CITY: Los Angeles
; STATE: California
; COUNTRY: USA
; ZIP: 90067
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
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;
; APPLICATION NUMBER: US/09/420.433
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/08/181,664
; FILING DATE: JANUARY 14, 1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Wetherell, Jr., Ph.D., John R.
; REGISTRATION NUMBER: 31,678
; REFERENCE/DOCKET NUMBER: PD-3055
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (619) 455-5100
; TELEFAX: (619) 455-5110
; INFORMATION FOR SEQ ID NO: 62:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 18 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: DNA (genomic)
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 1..18
US-09-420-433-62

Query Match 61.0%; Score 12.2; DB 10; Length 18;
Best Local Similarity 82.4%; Pred. No. 1.7e+03;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 TGCCTGACGTCAGAG 20
Db 1 TGCCTGCTGGGAG 17

RESULT 10
US-09-784-911-30/c
; Sequence 30, Application US/09784911
; Patent No. US20020072115A1
; GENERAL INFORMATION:
; APPLICANT: Harrison, Leonard C.
; APPLICANT: Jiang, Fang-Xu
; APPLICANT: Stanley, Edouard Guy
; APPLICANT: Genez, Leonel Jorge
; TITLE OF INVENTION: Pancreatic islet cell growth factors
; FILE REFERENCE: Davies Collison Cave
; CURRENT APPLICATION NUMBER: US/09/784,911
; CURRENT FILING DATE: 2001-09-17
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: PatentIn version 2.1
; SEQ ID NO 30
; LENGTH: 24
; TYPE: DNA
; ORGANISM: primer
US-09-784-911-30

Query Match 61.0%; Score 12.2; DB 10; Length 24;
Best Local Similarity 82.4%; Pred. No. 1.8e+03;
Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 4 TGCCTGACGTCAGAG 20
Db 17 TGCCTGACATCAAGAG 1

RESULT 11
US-09-769-066-11
; Sequence 11, Application US/09769066
; Patent No. US20020107360A1
; GENERAL INFORMATION:
; APPLICANT: Fuerst, Thomas R.
; McAttee, C. Patrick
; Yarbrough, Patrice O.
; Zhang, Yifan
```

;; TITLE OF INVENTION: HEPATITIS E VIRUS ANTIGENS AND USES THEREFOR  
;; NUMBER OF SEQUENCES: 31  
;; CORRESPONDENCE ADDRESS:

;; ADDRESSEE: Dehlinger & Associates  
;; STREET: 350 Cambridge Ave., Suite 250  
;; CITY: Palo Alto  
;; STATE: CA  
;; COUNTRY: USA

;; ZIP: 94306

;; COMPUTER READABLE FORM:

;; MEDIUM TYPE: Floppy disk

;; COMPUTER: IBM PC compatible

;; OPERATING SYSTEM: PC-DOS/MS-DOS

;; SOFTWARE: Patent In Release #1.0, Version #1.25

;; CURRENT APPLICATION DATA:

;; APPLICATION NUMBER: US/09/769,066

;; FILING DATE: 24-Jan-2001

;; CLASSIFICATION: <Unknown>

;; PRIOR APPLICATION DATA:

;; APPLICATION NUMBER: 08/542,634

;; FILING DATE: <Unknown>

;; ATTORNEY/AGENT INFORMATION:

;; NAME: Fabian, Gary R.

;; REGISTRATION NUMBER: 33,875

;; REFERENCE/DOCKET NUMBER: 4600-0293.30

;; TELECOMMUNICATION INFORMATION:

;; TELEPHONE: (415) 324-0880

;; TELEFAX: (415) 324-0960

;; INFORMATION FOR SEQ ID NO: 11:

;; SEQUENCE CHARACTERISTICS:

;; LENGTH: 99 base pairs

;; TYPE: nucleic acid

;; STRANDEDNESS: Hepatitis E Virus (Burma strain)

;; 406.4-2 region

;; TOPOLOGY: linear

;; MOLECULE TYPE: DNA (genomic)

;; HYPOTHETICAL: NO

;; ORIGINAL SOURCE:

;; SEQUENCE DESCRIPTION: SEQ ID NO: 11:

US-09-769-066-11

Query Match 61.0%; Score 12.2; DB 10; Length 99;

Best Local Similarity 82.4%; Pred. No. 2e+03;

Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 3 TTGCCTGACGTCAGAGA 19

||||| ||||| |||

DB 58 TTGCCTGACGTCGTAGA 74

RESULT 12

US-09-998-598-2339

;; Sequence 2339, Application US/09998598

;; Patent No. US20020150922A1

;; GENERAL INFORMATION:

;; APPLICANT: Stolk, John A.

;; APPLICANT: Xu, Jiangchun

;; APPLICANT: Chenault, Ruth A.

;; APPLICANT: Meagher, Madelein Joy

;; TITLE OF INVENTION: COMPOSITIONS AND METHODS FOR THE THERAPY AND

;; FILE REFERENCE: 210121.561

;; CURRENT APPLICATION NUMBER: US/09/998,598

;; NUMBER OF SEQ ID NOS: 2606

;; SOFTWARE: Corixa Invention Disclosure Database

;; SEQ ID NO 2339

;; LENGTH: 63

;; TYPE: DNA

;; ORGANISM: Homo sapiens

US-09-998-598-2339

Query Match 60.0%; Score 12; DB 10; Length 63;

Best Local Similarity 75.0%; Pred. No. 2.4e+03;  
Matches 15; Conservative 0; Mismatches 5; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20

||| ||||| ||||| |||

DB 14 GAACTCTGACGTCAGAG 33

RESULT 13

US-10-040-497-79

;; Sequence 79, Application US/10040497

;; Patent No. US20020172962A1

;; GENERAL INFORMATION:

;; APPLICANT: GOLD, LARRY

;; TUREK, CRAIG

;; TITLE OF INVENTION: METHODS OF PRODUCING NUCLEIC ACID LIGANDS

;; NUMBER OF SEQUENCES: 83

;; CORRESPONDENCE ADDRESS:

;; ADDRESSEE: Swanson & Bratschun, L.L.C.

;; STREET: 8400 E. Prentice Avenue, Suite 200

;; CITY: Englewood

;; STATE: Colorado

;; COUNTRY: USA

;; ZIP: 80111

;; COMPUTER READABLE FORM:

;; MEDIUM TYPE: Diskette, 3 1/5 inch, 1.44 MB

;; COMPUTER: IBM compatible

;; OPERATING SYSTEM: MS-DOS

;; SOFTWARE: WordPerfect 8.0

;; CURRENT APPLICATION DATA:

;; APPLICATION NUMBER: US/10/040,497

;; FILING DATE: 07-Jan-2002

;; PRIOR APPLICATION DATA:

;; APPLICATION NUMBER: 08/748,697

;; FILING DATE: 13-NOVEMBER-1996

;; APPLICATION NUMBER: 08/442,062

;; FILING DATE: 16-MAY-1995

;; APPLICATION NUMBER: 07/964,624

;; FILING DATE: 21-OCTOBER-1992

;; APPLICATION NUMBER: 07/714,131

;; FILING DATE: 10-JUNE-1991

;; APPLICATION NUMBER: 07/536,428

;; FILING DATE: 11-JUNE-1990

;; ATTORNEY/AGENT INFORMATION:

;; NAME: Barry J. Swanson

;; REGISTRATION NUMBER: 33,215

;; REFERENCE/DOCKET NUMBER: NEX05/DC-CON

;; TELECOMMUNICATION INFORMATION:

;; TELEPHONE: (303) 793-3333

;; TELEFAX: (303) 793-3433

;; INFORMATION FOR SEQ ID NO: 79:

;; SEQUENCE CHARACTERISTICS:

;; LENGTH: 86 base pairs

;; TYPE: nucleic acid

;; STRANDEDNESS: single

;; TOPOLOGY: linear

;; SEQUENCE DESCRIPTION: SEQ ID NO: 79:

US-10-040-497-79

Query Match

Best Local Similarity 60.0%; Score 12; DB 9; Length 86;

Matches 13; Conservative 2; Mismatches 5; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAG 20

||| ||||| ||||| |||

DB 10 GAUGGCCUCCGACCGAG 29

RESULT 14

US-09-864-761-28968

;; Sequence 28968, Application US/09864761

;; Patent No. US20020048763A1

;; GENERAL INFORMATION:

```

RESULT 15
US-08-983-605-140
; Sequence 140, Application US/08983605A
; Patent No. US20020066118A1
; GENERAL INFORMATION:
; APPLICANT: Roder, Marion
; TITLE OF INVENTION: Microsatellite Markers for plants of the species
; TITLE OF INVENTION: Triticum aestivum and Tribe Triticaceae and the Use of
; TITLE OF INVENTION: Said Markers
; FILE REFERENCE: 2936.10400
; CURRENT APPLICATION NUMBER: US/08/983.605A
; CURRENT FILING DATE: 1998-05-01
; EARLIER APPLICATION NUMBER: DE 195 25 284.5
; EARLIER FILING DATE: 1995-06-28
; NUMBER OF SEQ ID NOS: 466
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 140
; LENGTH: 18
; TYPE: DNA
; ORGANISM: Triticum aestivum
US-08-983-605-140

Query Match          59.0%;   Score 11.8;   DB 8;   Length 18;
Best Local Similarity 86.7%;
Matches 13;   Conservative    0;   Mismatches    2;   Indels    0;   Gaps

QY      6  CCTGACGTCAGAGAG 20
        ||| ||||| |||||
DB      4  CCCACGTCAGAGAG 18

Search completed: December 12, 2002, 06:12:14
Job time : 42.0549 secs

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Query Match 60.0%; Score 12; DB 10; Length 92;  
Best Local Similarity 75.0%; Pred. No. 2.5e+03;  
Matches 15; Conservative 0; Mismatches 5; Indels

GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:05:40 ; Search time 22.2464 Seconds  
(without alignments)  
275.709 Million cell updates/sec

Title: US-09-355-254F-8  
Perfect score: 20  
Sequence: 1 gattgctgacgtcagagag 20

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 153338381 residues

Total number of hits satisfying chosen parameters: 687286

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : Issued\_Patents\_NA.\*  
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2: /cgn2\_6/ptodata/1/ina/5B\_COMB.seq.\*  
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4: /cgn2\_6/ptodata/1/ina/6B\_COMB.seq.\*  
5: /cgn2\_6/ptodata/1/ina/PCTUS\_COMB.seq.\*  
6: /cgn2\_6/ptodata/1/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	US-09-133-774-10	Sequence 10, Appl
2	20	100.0	20	US-09-303-862-10	Sequence 10, Appl
3	20	100.0	27	US-08-210-880B-2	Sequence 2, Appl
4	20	100.0	27	US-08-771-411-2	Sequence 2, Appl
5	16.4	82.0	32	US-09-215-098-1	Sequence 1, Appl
6	16.4	82.0	50	US-08-171-389-451	Sequence 451, App
7	16.4	82.0	50	US-08-123-936-451	Sequence 451, App
8	16.4	82.0	50	US-08-475-228A-451	Sequence 451, App
9	16.4	82.0	50	US-08-482-080A-451	Sequence 451, App
10	16.4	82.0	50	US-09-354-947-451	Sequence 451, App
11	16.4	82.0	50	PCT-US93-12388-451	Sequence 451, App
12	14.2	71.0	41	US-08-813-507-78	Sequence 78, Appl
13	14.2	71.0	41	US-09-464-453-78	Sequence 78, Appl
14	13.8	69.0	71	US-08-870-930-25	Sequence 25, Appl
15	13.8	69.0	71	US-09-254-968-28	Sequence 28, Appl
16	12.8	64.0	61	US-09-023-228B-89	Sequence 89, Appl
17	12.8	64.0	61	US-09-163-025B-89	Sequence 89, Appl
18	12.6	63.0	57	US-08-701-124-60	Sequence 60, Appl
19	12.6	63.0	57	US-08-701-124-61	Sequence 61, Appl
20	12.6	63.0	57	US-09-130-225-60	Sequence 60, Appl
21	12.6	63.0	57	US-09-130-225-61	Sequence 61, Appl
22	12.6	63.0	57	US-09-455-061-60	Sequence 60, Appl
23	12.6	63.0	57	US-09-455-061-61	Sequence 61, Appl
24	12.6	63.0	63	US-08-701-124-58	Sequence 58, Appl
25	12.6	63.0	63	US-09-130-225-58	Sequence 58, Appl
26	12.6	63.0	63	US-09-455-061-58	Sequence 58, Appl
27	12.6	63.0	98	US-08-441-430-5	Sequence 5, Appl

c	28	12.4	62.0	27	2	US-08-244-434-13	Sequence 13, Appl
	29	12.4	62.0	35	2	US-08-244-434-12	Sequence 12, Appl
	30	12.4	62.0	35	2	US-08-244-434-16	Sequence 16, Appl
c	31	12.4	62.0	40	2	US-09-130-114-18	Sequence 18, Appl
	32	12.4	62.0	44	2	US-09-130-114-17	Sequence 17, Appl
	33	12.4	62.0	18	1	US-08-152-313-89	Sequence 89, Appl
	34	12.2	61.0	18	1	US-08-579-223-89	Sequence 89, Appl
	35	12.2	61.0	18	3	US-08-181-664-62	Sequence 62, Appl
	36	12.2	61.0	18	5	PCT-US94-12947A-89	Sequence 89, Appl
c	37	12.2	61.0	20	4	US-09-122-171D-15	Sequence 15, Appl
	38	12.2	61.0	21	4	US-09-324-867-43	Sequence 43, Appl
c	39	12.2	61.0	22	3	US-08-840-316-95	Sequence 95, Appl
	40	12.2	61.0	22	4	US-08-809-523-95	Sequence 95, Appl
c	41	12.2	61.0	22	4	US-08-471-971-95	Sequence 95, Appl
	42	12.2	61.0	22	4	US-09-402-776-95	Sequence 95, Appl
c	43	12.2	61.0	22	5	PCT-US93-08849A-95	Sequence 95, Appl
	44	12.2	61.0	22	5	PCT-US93-08849-95	Sequence 95, Appl
c	45	12.2	61.0	25	1	US-07-991-466-1	Sequence 1, Appl

ALIGNMENTS

RESULT 1  
US-09-133-774-10  
; Sequence 10, Application US/09133774B  
; Patent No. 5962636  
; GENERAL INFORMATION:  
; APPLICANT: Bachmaier, Kurt  
; APPLICANT: Hessel, Andrew J.  
; APPLICANT: Neu M.D., Nikolaus  
; APPLICANT: Penninger, Josef M.  
; TITLE OF INVENTION: No. 5962636el Peptides Capable of Modulating Inflammatory Hear  
; TITLE OF INVENTION: Disease  
; FILE REFERENCE: A-536  
; CURRENT APPLICATION NUMBER: US/09/133,774B  
; CURRENT FILING DATE: 1998-08-12  
; NUMBER OF SEQ ID NOS: 26  
; SOFTWARE: Patentin Ver. 2.0  
; SEQ ID NO 10  
; LENGTH: 20  
; TYPE: DNA  
; ORGANISM: Chlamydia trachomatis  
; FEATURE:  
; OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a  
; OTHER INFORMATION: 60 kDa cysteine rich outer membrane protein from  
; OTHER INFORMATION: Chlamydia trachomatis.  
US-09-133-774-10

Query Match 100.0%; Score 20; DB 2; Length 20;  
Best Local Similarity 100.0%; Pred No. 0.057; 0; Gaps 0;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GATTGCTGACGTGACAGAG 20  
Db 1 GATTGCTGACGTGACAGAG 20  
|||||

RESULT 2  
US-09-303-862-10  
; Sequence 10, Application US/09303862  
; Patent No. 6034230  
; GENERAL INFORMATION:  
; APPLICANT: Bachmaier, Kurt  
; APPLICANT: Hessel, Andrew J.  
; APPLICANT: Neu M.D., Nikolaus  
; APPLICANT: Penninger, Josef M.  
; TITLE OF INVENTION: No. 6034230el Peptides Capable of Modulating Inflammatory Hear  
; TITLE OF INVENTION: Disease  
; FILE REFERENCE: A-536  
; CURRENT APPLICATION NUMBER: US/09/303,862  
; CURRENT FILING DATE: 1999-05-03  
; EARLIER APPLICATION NUMBER: 09/133,774

EARLIER FILING DATE: 1998-08-12  
NUMBER OF SEQ ID NOS: 26  
SOFTWARE: PatentIn Ver. 2.0  
SEQ ID NO 10  
LENGTH: 20  
TYPE: DNA  
ORGANISM: Chlamydia trachomatis  
FEATURE:  
OTHER INFORMATION: An oligonucleotide derived from the DNA encoding a  
OTHER INFORMATION: 60 kDa cysteine rich outer membrane protein from  
OTHER INFORMATION: Chlamydia trachomatis.  
US-09-303-862-10

Query Match 100.0%; Score 20; DB 3; Length 20;  
Best Local Similarity 100.0%; Pred. No. 0.057;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GATTGCTGACGTCAGAG 20  
Db 1 GATTGCTGACGTCAGAG 20  
|||||

RESULT 3  
US-08-210-880B-2  
Sequence 2, Application US/08210880B  
Patent No. 5641486  
GENERAL INFORMATION:  
APPLICANT: HINRICHS, STEVEN H.  
APPLICANT: ORTEN, DANA J.  
TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING  
NUMBER OF INVENTION: CREB/ATF1 FAMILY DETERMINED TRANSCRIPTION  
NUMBER OF SEQUENCES: 11  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: HENDERSON & STURM  
STREET: 1125 S. 103RD ST., #330  
CITY: OMAHA  
STATE: NE  
COUNTRY: US  
ZIP: 68124  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/210,880B  
FILING DATE: 18-MAR-1994  
CLASSIFICATION: 424  
ATTORNEY/AGENT INFORMATION:  
NAME: JONDLE, ROBERT J.  
REGISTRATION NUMBER: 33,915  
REFERENCE/DOCKET NUMBER: 63066  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 402-398-9000  
TELEFAX: 402-398-9005  
INFORMATION FOR SEQ ID NO: 2:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 27 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: unknown  
MOLECULE TYPE: DNA (genomic)  
US-08-210-880B-2

Query Match 100.0%; Score 20; DB 1; Length 27;  
Best Local Similarity 100.0%; Pred. No. 0.059;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GATTGCTGACGTCAGAG 20  
Db 4 GATTGCTGACGTCAGAG 23  
|||||

RESULT 4  
US-08-771-411-2  
Sequence 2, Application US/08771411  
Patent No. 5844096  
GENERAL INFORMATION:  
APPLICANT: HINRICHS, STEVEN H.  
APPLICANT: ORTEN, DANA J.  
TITLE OF INVENTION: METHOD AND COMPOSITION FOR INHIBITING  
NUMBER OF INVENTION: CREB/ATF1 FAMILY DETERMINED TRANSCRIPTION  
NUMBER OF SEQUENCES: 11  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: HENDERSON & STURM  
STREET: 1125 S. 103RD ST., #330  
CITY: OMAHA  
STATE: NE  
COUNTRY: US  
ZIP: 68124  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/771,411  
FILING DATE: 20-DEC-1996  
CLASSIFICATION: 424  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/210,880  
FILING DATE: 18-MAR-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: JONDLE, ROBERT J.  
REGISTRATION NUMBER: 33,915  
REFERENCE/DOCKET NUMBER: 63066  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 402-398-9000  
TELEFAX: 402-398-9005  
INFORMATION FOR SEQ ID NO: 2:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 27 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: unknown  
MOLECULE TYPE: DNA (genomic)  
US-08-771-411-2

Query Match 100.0%; Score 20; DB 2; Length 27;  
Best Local Similarity 100.0%; Pred. No. 0.059;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

Qy 1 GATTGCTGACGTCAGAG 20  
Db 4 GATTGCTGACGTCAGAG 23  
|||||

RESULT 5  
US-09-215-098-1  
Sequence 1, Application US/09215098  
Patent No. 6194632  
GENERAL INFORMATION:  
APPLICANT: Leiden, Jeffery M  
TITLE OF INVENTION: DILATED CARDIOMYOPATHY IN TRANSGENIC MICE EXPRESSING A  
TITLE OF INVENTION: DOMINANT-NEGATIVE CREB TRANSCRIPTION FACTOR IN THE  
TITLE OF INVENTION: HEART  
FILE REFERENCE: 9189-4  
CURRENT APPLICATION NUMBER: US/09/215,098  
CURRENT FILING DATE: 1998-12-18  
PRIOR APPLICATION NUMBER: 60/068,011  
PRIOR FILING DATE: 1997-12-18  
NUMBER OF SEQ ID NOS: 5  
SOFTWARE: PatentIn Ver. 2.1  
SEQ ID NO 1  
LENGTH: 32  
TYPE: DNA



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RESULT 8
US-08-475-228A-451
; Sequence 451, Application US/08475228A
; Patent No. 5869241
; GENERAL INFORMATION:
; APPLICANT: Edwards, Cynthia A.
; APPLICANT: Cantor, Charles R.
; APPLICANT: Andrews, Beth M.
; APPLICANT: Turin, Lisa M.
; APPLICANT: Fry, Kirk E.
; TITLE OF INVENTION: Sequence-Directed DNA Binding
; MOLECULES, COMPOSITIONS AND METHODS
; NUMBER OF SEQUENCES: 664
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Genelabs Technologies, Inc.

```

STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: Patentin Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/475,228A  
FILING DATE: 06-JUN-1995  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
PRIORITY APPLICATION DATA:  
APPLICATION NUMBER: US 08/081,070  
FILING DATE: 22-JUN-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Stratford Carol A.  
REGISTRATION NUMBER: 34,444  
REFERENCE/DOCKET NUMBER: 4600-0175.21/G19P3D2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 324-0880  
TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 451:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 50 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Human somatostatin I gene  
US-08-475-228A-451

Query Match	82.0%	Score 15.4;	DB 2;	Length 50;
Best Local Similarity	94.4%	Pred. No. 5.2;		
Matches 17;	Conservative	0;	Mismatches 1;	Indels
QY	3	TTGCCTGACGTCAGAG	20	
Db	1	TAGCCTGACGTCAGAG	18	

RESULT 9  
US-08-482-080A-451  
; Sequence 451, Application US/08482080A  
; Patent No. 6010849.

```

: GENERAL INFORMATION:
: APPLICANT: Edwards, Cynthia A.
: APPLICANT: Cantor, Charles R.
: APPLICANT: Andrews, Beth M.
: APPLICANT: Turin, Lisa M.
: APPLICANT: Fry, Kirk E.
: TITLE OF INVENTION: Sequence-Directed DNA Binding
: TITLE OF INVENTION: Molecules, Compositions and Methods
: NUMBER OF SEQUENCES: 664
: CORRESPONDENCE ADDRESS:
: ADDRESSEE: Genelabs Technologies, Inc.
: STREET: 505 Penobscot Drive
: CITY: Redwood City
: STATE: CA
: COUNTRY: USA
: ZIP: 94063
: COMPUTER READABLE FORM:
: MEDIUM TYPE: Floppy disk
: COMPUTER: IBM PC compatible
: OPERATING SYSTEM: PC-DOS/MS-DOS
: SOFTWARE: Patentin Release #1.0, Version #1.25
: CURRENT APPLICATION DATA:
: APPLICATION NUMBER: US/08/482.080A
: FILING DATE: 07-JUN-1995
: PRIORITY APPLICATION DATA:
: APPLICATION NUMBER: US 08/171,389
: FILING DATE: 20-DEC-1993
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/123,936
: FILING DATE: 17-SEP-1993
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/996,783
: FILING DATE: 23-DEC-1992
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 07/723,618
: FILING DATE: 27-JUN-1991
: PRIOR APPLICATION DATA:
: APPLICATION NUMBER: US 08/081,070
: FILING DATE: 22-JUN-1993
: ATTORNEY/AGENT INFORMATION:
: NAME: Brady, John F.
: REGISTRATION NUMBER: 39,118
: REFERENCE/DOCKET NUMBER: 4600-0175.20/G19p3D1
: TELECOMMUNICATION INFORMATION:
: TELEPHONE: (650) 324-0880
: TELEFAX: (650) 324-0960
: INFORMATION FOR SEQ ID NO: 451:
: SEQUENCE CHARACTERISTICS:
: LENGTH: 50 base pairs
: TYPE: nucleic acid
: STRANDEDNESS: double
: TOPOLOGY: linear
: MOLECULE TYPE: DNA (genomic)
: HYPOTHEetical: NO
: ORIGINAL SOURCE:
: INDIVIDUAL ISOLATE: Human somatostatin 1 gene
: US-08-482-080A-451

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Query Match	82.08;	Score 16.4;	DB 3;	Length 50;
Best Local Similarity	94.4;	Pred. No. 5.2;		
Matches 17; Conservative	0;	Mismatches	1;	Indels 0; Gaps 0;
Qy	3	TTGCTGACGCTCAGAG	20	
Db	1	TAGCTGACGCTCAGAG	18	

RESULT 10  
US-09-354-947-451  
; Sequence 451, Application US/09354947  
; Patent No. 6384208  
; GENERAL INFORMATION:  
; APPLICANT: Edwards, Cynthia A.





SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/813,507  
FILING DATE: 07-MAR-1997  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 60/032,069  
FILING DATE: 02-DEC-1996  
ATTORNEY/AGENT INFORMATION:  
NAME: Liebeschuetz, Joe  
REGISTRATION NUMBER: 37,505  
REFERENCE/DOCKET NUMBER: 018547-030100US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415 576-0200  
TELEFAX: 415 576-0200  
TELEX:  
INFORMATION FOR SEQ ID NO: 78:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 41 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
US-08-813-507-78

Query Match 71.0%; Score 14.2; DB 3; Length 41;  
Best Local Similarity 84.2%; Pred. No. 74;  
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAGA 19  
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DB 19 GCTTTCATGACGTCAGAGA 1

## RESULT 13

US-09-464-453-78/c  
Sequence 78, Application US/09464453  
Patent No. 6358686  
GENERAL INFORMATION:  
APPLICANT: Lemieux, Bertrand  
Sepolsky, Ronald J.  
TITLE OF INVENTION: Brassinica Polymorphisms  
NUMBER OF SEQUENCES: 173  
CORRESPONDENCE ADDRESS:  
ADDRESS: Townsend and Townsend and Crew LLP  
STREET: Two Embarcadero Center, Eighth Floor  
CITY: San Francisco  
STATE: California  
COUNTRY: USA  
ZIP: 94111-3834  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette  
COMPUTER: IBM Compatible  
OPERATING SYSTEM: DOS  
SOFTWARE: FastSeq for Windows Version 2.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/464,453  
FILING DATE: 14-DEC-1999  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: 08/813,507  
FILING DATE: <Unknown>  
ATTORNEY/AGENT INFORMATION:  
NAME: Liebeschuetz, Joe  
REGISTRATION NUMBER: 37,505  
REFERENCE/DOCKET NUMBER: 018547-030100US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: 415 576-0200  
TELEFAX: 415 576-0200  
TELEX: <Unknown>  
INFORMATION FOR SEQ ID NO: 78:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 41 base pairs

TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
SEQUENCE DESCRIPTION: SEQ ID NO: 78:  
US-09-464-453-78

Query Match 71.0%; Score 14.2; DB 4; Length 41;  
Best Local Similarity 84.2%; Pred. No. 74;  
Matches 16; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAGA 19  
| | | | | | | | | | |  
DB 19 GCTTTCATGACGTCAGAGA 1

## RESULT 14

US-08-870-930-25  
Sequence 25, Application US/08870930  
Patent No. 6168778  
GENERAL INFORMATION:  
APPLICANT: NEBOJSA JANJIC, LARRY GOLD, PAUL G. SCHMIDT, CHANDRA VARGESE, MICHA  
TITLE OF INVENTION: VASCULAR ENDOTHELIAL GROWTH FACTOR (VEGF)  
NUMBER OF SEQUENCES: 88  
CORRESPONDENCE ADDRESS:  
ADDRESS: Swanson and Bratschun, L.L.C.  
STREET: 8400 East Prentice Avenue, Suite #200  
CITY: Denver  
STATE: Colorado  
COUNTRY: USA  
ZIP: 80111

COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.5 inch, 1.4 Mb storage  
COMPUTER: IBM compatible  
OPERATING SYSTEM: MS-DOS  
SOFTWARE: WordPerfect 8.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/870,930  
FILING DATE: 6 JUNE 1997  
CLASSIFICATION: 536

ATTORNEY/AGENT INFORMATION:  
NAME: Barry J. Swanson  
REGISTRATION NUMBER: 33,215  
REFERENCE/DOCKET NUMBER: NEX61  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (303) 793-3333  
TELEFAX: (303) 793-3433  
INFORMATION FOR SEQ ID NO: 25:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 71  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: RNA  
FEATURE:  
OTHER INFORMATION: All pyrimidines are 2'-fluoro

US-08-870-930-25

Query Match 69.0%; Score 13.8; DB 4; Length 71;  
Best Local Similarity 64.7%; Pred. No. 1.3e+02;  
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGA 17  
| | | | | | | | | | |  
DB 43 GAUUUCCGCGCUCAGA 59

## RESULT 15

US-09-254-968-28  
Sequence 28, Application US/09254968  
Patent No. 6426335  
GENERAL INFORMATION:  
APPLICANT: NEBOJSA JANJIC, LARRY GOLD, PAUL G. SCHMIDT, CHANDRA VARGESE,

MICHAEL WILLIS  
TITLE OF INVENTION: VASCULAR ENDOTHELIAL GROWTH FACTOR (VEGF) NUCLEIC  
ACID LIGAND COMPLEXES  
NUMBER OF SEQUENCES: 139  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Swanson and Bratschun, L.L.C.  
STREET: 8400 East Prentice Avenue, Suite #200  
CITY: Denver  
STATE: Colorado  
COUNTRY: USA  
ZIP: 80111  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Diskette, 3.5 inch, 1.4 Mb storage  
COMPUTER: IBM compatible  
(C) OPERATING SYSTEM: MS-DOS  
SOFTWARE: Word 7.0  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/254,968  
FILING DATE: 13-Mar-2000  
CLASSIFICATION: <Unknown>  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: PCT/US97/18944  
FILING DATE: 17 OCTOBER 1997  
APPLICATION NUMBER: 08/739,109  
FILING DATE: 25 OCTOBER 1996  
APPLICATION NUMBER: 08/870,930  
FILING DATE: 6 JUNE 1997  
APPLICATION NUMBER: 08/897,351  
FILING DATE: 21 JULY 1997  
ATTORNEY/AGENT INFORMATION:  
NAME: Barry J. Swanson  
REGISTRATION NUMBER: 33,215  
REFERENCE/DOCKET NUMBER: NEX61C/PCT-US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (303) 793-3333  
TELEFAX: (303) 793-3433  
INFORMATION FOR SEQ ID NO: 28:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 71  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear  
MOLECULE TYPE: RNA  
FEATURE:  
OTHER INFORMATION: All pyrimidines are 2'-fluoro (2'-F)  
modified  
SEQUENCE DESCRIPTION: SEQ ID NO: 28:  
US-09-254-968-28  
Query Match 69.0%; Score 13.8; DB 4; Length 71;  
Best Local Similarity 64.7%; Pred. No. 1.3e+02;  
Matches 11; Conservative 4; Mismatches 2; Indels 0; Gaps 0;  
QY 1 GATTGCTGACGTCAGA 17  
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Db 43 GAUUUCCUGCCGUCAGA 59

Search completed: December 12, 2002, 01:41:42  
Job time : 24.2464 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds  
(without alignments)  
1829.698 Million cell updates/sec

Title: US-09-355-254F-13

Perfect score: 20  
Sequence: 1 tgcagattgcgaactgtca 20

Scoring table: IDENTITY\_NUC  
Gapop 10\_0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : GenEmbl.\*

- 1: gb\_ba.\*
- 2: gb\_htg.\*
- 3: gb\_in.\*
- 4: gb\_om.\*
- 5: gb\_ov.\*
- 6: gb\_pat.\*
- 7: gb\_ph.\*
- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
- 12: gb\_sy.\*
- 13: gb\_un.\*
- 14: gb\_vi.\*
- 15: em\_ba.\*
- 16: em\_fun.\*
- 17: em\_hum.\*
- 18: em\_in.\*
- 19: em\_mu.\*
- 20: em\_om.\*
- 21: em\_or.\*
- 22: em\_ov.\*
- 23: em\_pat.\*
- 24: em\_ph.\*
- 25: em\_pl.\*
- 26: em\_ro.\*
- 27: em\_sts.\*
- 28: em\_un.\*
- 29: em\_vi.\*
- 30: em\_htg\_hum.\*
- 31: em\_htg\_inv.\*
- 32: em\_htg\_other.\*
- 33: em\_htg\_mus.\*
- 34: em\_htg\_pln.\*
- 35: em\_htg\_rod.\*
- 36: em\_htg\_mam.\*
- 37: em\_htg\_vrt.\*
- 38: em\_sy.\*
- 39: em\_htgo\_hum.\*
- 40: em\_htgo\_mus.\*
- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	20	100.0	20	6	A89792	A89792 Sequence 14
2	20	100.0	20	6	A89792	A89792 Sequence 14
3	20	100.0	20	6	A90879	A90879 Sequence 14
4	20	100.0	20	6	A90879	A90879 Sequence 14
5	20	100.0	20	6	AR148641	AR148641 Sequence
6	20	100.0	20	6	AR148641	AR148641 Sequence
7	20	100.0	20	6	AX455640	AX455640 Sequence
8	20	100.0	20	6	AX455640	AX455640 Sequence
9	15.4	77.0	24	6	AX384713	AX384713 Sequence
10	15.4	77.0	24	6	AX384713	AX384713 Sequence
11	13.6	68.0	90	6	AR024336	AR024336 Sequence
12	13.6	68.0	90	6	AR024336	AR024336 Sequence
13	13.6	68.0	90	6	AR045189	AR045189 Sequence
14	13.6	68.0	90	6	AR045189	AR045189 Sequence
15	13.6	68.0	90	6	BD011413	BD011413 Chimeric
16	13.6	68.0	90	6	BD011413	BD011413 Chimeric
17	13.6	68.0	90	6	E43883	E43883 Chimeric an
18	13.6	68.0	90	6	E43883	E43883 Chimeric an
19	13.4	67.0	28	6	AR202106	AR202106 Sequence
20	13.4	67.0	28	6	AR202106	AR202106 Sequence
21	13.2	66.0	21	6	AR041200	AR041200 Sequence
22	13.2	66.0	21	6	AR041200	AR041200 Sequence
23	13.2	66.0	21	6	AR116609	AR116609 Sequence
24	13.2	66.0	21	6	AR116609	AR116609 Sequence
25	13.2	66.0	21	6	AR159952	AR159952 Sequence
26	13.2	66.0	21	6	AR159952	AR159952 Sequence
27	13.2	66.0	21	6	AX268106	AX268106 Sequence
28	13.2	66.0	21	6	AX268106	AX268106 Sequence
29	13.2	66.0	33	6	I65281	I65281 Sequence 3
30	13.2	66.0	33	6	I65281	I65281 Sequence 3
31	13.2	66.0	54	6	AX244174	AX244174 Sequence
32	13.2	66.0	54	6	AX244174	AX244174 Sequence
33	13.2	66.0	59	6	AX015212	AX015212 Sequence
34	13.2	66.0	59	6	AX015212	AX015212 Sequence
35	13.2	66.0	59	6	E64355	E64355 Single-stra
36	13.2	66.0	59	6	E64355	E64355 Single-stra
37	13.2	66.0	90	6	AX287868	AX287868 Sequence
38	13.2	66.0	90	6	AX287868	AX287868 Sequence
39	12.8	64.0	56	6	A62184	A62184 Sequence 78
40	12.8	64.0	56	6	A62184	A62184 Sequence 78
41	12.8	64.0	56	6	AR077589	AR077589 Sequence
42	12.8	64.0	56	6	AR077589	AR077589 Sequence
43	12.8	64.0	56	6	AX022408	AX022408 Sequence
44	12.8	64.0	56	6	AX022408	AX022408 Sequence
45	12.8	64.0	56	6	AX047755	AX047755 Sequence

ALIGNMENTS

RESULT 1  
A89792  
LOCUS A89792  
DEFINITION Sequence 14 from Patent WO9832462.  
ACCESSION A89792  
VERSION A89792.1 GI:6738306  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unclassified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Lipford,G.B. and Heeg,K.  
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
JOURNAL Patent: WO 9832462-A 14 30-JUL-1998;

linear PAT 22-JAN-2000

-AUG-2001

ACCESSION ARI48641  
VERSION ARI48641.1 GI:15112731  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Shashoua, V.E.  
TITLE Neuroprotective peptides and uses thereof  
JOURNAL Patent: US 6225444-A 7 01-MAY-2001;  
FEATURES Location/Qualifiers  
source 1..20  
BASE COUNT 5 a 5 c 5 g 5 t  
ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;  
Best Local Similarity 100.0%; Pred. No. 22;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCAGATTGCGCAATCTGCA 20  
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Db 20 TGCAGATTGCGCAATCTGCA 1

RESULT 7  
LOCUS AX455640 20 bp DNA linear PAT 06-JUL-2002  
DEFINITION Sequence 117 from Patent WO0222809.  
ACCESSION AX455640  
VERSION AX455640.1 GI:21714708  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1  
AUTHORS Bauer, S., Lipford, G. and Wagner, H.  
TITLE Process for high throughput screening of cpq-based  
JOURNAL immuno-agonist/antagonist  
Patent: WO 0222809-A 117 21-MAR-2002;  
Coley Pharmaceutical GmbH (DE)  
FEATURES Location/Qualifiers  
source 1..20  
BASE COUNT 5 a 5 c 5 g 5 t  
ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;  
Best Local Similarity 100.0%; Pred. No. 22;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCAGATTGCGCAATCTGCA 20  
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Db 1 TGCAGATTGCGCAATCTGCA 20

RESULT 8  
LOCUS AX455640/c 20 bp DNA linear PAT 06-JUL-2002  
DEFINITION Sequence 117 from Patent WO0222809.  
ACCESSION AX455640  
VERSION AX455640.1 GI:21714708  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1  
AUTHORS Bauer, S., Lipford, G. and Wagner, H.  
TITLE Process for high throughput screening of cpq-based  
JOURNAL immuno-agonist/antagonist  
Patent: WO 0222809-A 117 21-MAR-2002;

ACCESSION ARI48641  
VERSION ARI48641.1 GI:15112731  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Shashoua, V.E.  
TITLE Neuroprotective peptides and uses thereof  
JOURNAL Patent: US 6225444-A 7 01-MAY-2001;  
FEATURES Location/Qualifiers  
source 1..20  
BASE COUNT 5 a 5 c 5 g 5 t  
ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 20;  
Best Local Similarity 100.0%; Pred. No. 22;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TGCAGATTGCGCAATCTGCA 20  
|||||  
Db 20 TGCAGATTGCGCAATCTGCA 1

RESULT 9  
LOCUS AX384713 24 bp DNA linear PAT 19-MAR-2002  
DEFINITION Sequence 4 from Patent WO0181375.  
ACCESSION AX384713  
VERSION AX384713.1 GI:19577904  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1  
AUTHORS Schepartz Shrader, A., Chin, J.W., Zutshi, R., Rutledge, S.E., Kehlbeck  
TITLE Dna & protein binding miniature proteins  
JOURNAL Patent: WO 0181375-A 4 01-NOV-2001;  
YALE UNIVERSITY (US)  
FEATURES Location/Qualifiers  
source 1..24  
BASE COUNT 5 a 5 c 8 g 6 t  
ORIGIN

Query Match 77.0%; Score 15.4; DB 6; Length 24;  
Best Local Similarity 94.1%; Pred. No. 5.5e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 TCCAGATTGCGCAATCT 17  
|||||  
Db 3 TGGAGATTGCGCAATCT 19

RESULT 10  
LOCUS AX384713/c 24 bp DNA linear PAT 19-MAR-2002  
DEFINITION Sequence 4 from Patent WO0181375.  
ACCESSION AX384713  
VERSION AX384713.1 GI:19577904  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1  
AUTHORS Schepartz Shrader, A., Chin, J.W., Zutshi, R., Rutledge, S.E., Kehlbeck  
TITLE Dna & protein binding miniature proteins  
JOURNAL Patent: WO 0181375-A 4 01-NOV-2001;  
YALE UNIVERSITY (US)  
FEATURES Location/Qualifiers  
source 1..24  
BASE COUNT 5 a 5 c 8 g 6 t  
ORIGIN

ORIGIN

Query Match 77.0%; Score 15.4; DB 6; Length 24;  
Best Local Similarity 94.18; Pred. No. 5.5e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 4 AGATTGCGCAATCTGCA 20  
||||| ||||| ||||| ||  
Db 19 AGATTGCGCAATCTGCA 3

RESULT 11  
AR024336  
LOCUS AR024336 90 bp DNA linear PAT 05-DEC-1998  
DEFINITION Sequence 104 from patent US 5795965.  
ACCESSION AR024336  
VERSION AR024336.1 GI:3977630  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)  
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.  
TITLE Reshaped human antibody to human interleukin-6 receptor  
JOURNAL Patent: US 5795965-A 104 18-AUG-1998;  
FEATURES Location/Qualifiers  
source 1..90  
BASE COUNT 18 a 22 c 29 g 21 t  
ORIGIN

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Best Local Similarity 80.0%; Pred. No. 4.4e+04;  
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

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||||| ||||| ||||| ||  
Db 11 TGCAGCTTGTGCAGTCTGGA 30

RESULT 12  
AR024336/c  
LOCUS AR024336 90 bp DNA linear PAT 05-DEC-1998  
DEFINITION Sequence 104 from patent US 5795965.  
ACCESSION AR024336  
VERSION AR024336.1 GI:3977630  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)  
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.  
TITLE Reshaped human antibody to human interleukin-6 receptor  
JOURNAL Patent: US 5795965-A 104 18-AUG-1998;  
FEATURES Location/Qualifiers  
source 1..90  
BASE COUNT 18 a 22 c 29 g 21 t  
ORIGIN

Query Match 68.0%; Score 13.6; DB 6; Length 90;  
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Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 TGCAGATTGCGCAATCTGCA 20  
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Db 30 TCCAGACTGCACAGCTGCA 11

RESULT 13  
AR045189

LOCUS AR045189 90 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 104 from patent US 5817790.  
ACCESSION AR045189  
VERSION AR045189.1 GI:5966654  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)  
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.  
TITLE Reshaped human antibody to human interleukin-6 receptor  
JOURNAL Patent: US 5817790-A 104 06-OCT-1998;  
FEATURES Location/Qualifiers  
source 1..90  
BASE COUNT 18 a 22 c 29 g 21 t  
ORIGIN

Query Match 68.0%; Score 13.6; DB 6; Length 90;  
Best Local Similarity 80.0%; Pred. No. 4.4e+04;  
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 TGCAGATTGCGCAATCTGCA 20  
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Db 11 TGCAGCTTGTGCAGTCTGGA 30

RESULT 14  
AR045189/c  
LOCUS AR045189 90 bp DNA linear PAT 29-SEP-1999  
DEFINITION Sequence 104 from patent US 5817790.  
ACCESSION AR045189  
VERSION AR045189.1 GI:5966654  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.

REFERENCE 1 (bases 1 to 90)  
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.Margaret., Jones,S.Tarran. and Saldanha,J.William.  
TITLE Reshaped human antibody to human interleukin-6 receptor  
JOURNAL Patent: US 5817790-A 104 06-OCT-1998;  
FEATURES Location/Qualifiers  
source 1..90  
BASE COUNT 18 a 22 c 29 g 21 t  
ORIGIN

Query Match 68.0%; Score 13.6; DB 6; Length 90;  
Best Local Similarity 80.0%; Pred. No. 4.4e+04;  
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

Qy 1 TGCAGATTGCGCAATCTGCA 20  
||||| ||||| ||||| ||  
Db 30 TCCAGACTGCACAGCTGCA 11

RESULT 15  
BD011413  
LOCUS BD011413 90 bp DNA linear PAT 31-JAN-2002  
DEFINITION Chimeric antibody against human interleukin-6 receptor.  
ACCESSION BD011413  
VERSION BD011413.1 GI:18639786  
KEYWORDS JP 2001083151-A/87.  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 90)  
AUTHORS Tsuchiya,M., Sato,K., Bendig,M.M., Jones,S.T. and Saldanha,H.W.  
TITLE Chimeric antibody against human interleukin-6 receptor  
JOURNAL Patent: JP 2001083151-A 87 30-MAR-2001;  
CHUGAI PHARMACEUTICAL CO LTD



COMMENT OS- Artificial Sequence  
PN JP 2001083151-A/87  
PD 30-MAR-2001  
PF 28-JUL-2000 JP 2000229748  
PR  
PI MASAYUKI TSUCHIYA, KO SATO, MARY MARGARET BENDIGU, PI STEVEN  
TAREN JONES,  
PI HOSE WILLIAM SALDANHA  
PC G01N33/53, A61K38/00, A61K39/395, A61K39/395, A61P35/00, PC  
G01N33/577//C07K16/28,  
PC C07K19/00, C12N15/09, (C12N15/09, C12R1:91), A61K37/02, C12N15/00,  
PC (C12N15/00, C12R1:91)  
CC  
FH Key Location/Qualifiers  
FT source 1..90  
FT /organism='Artificial Sequence'.  
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1..90  
/organism='synthetic construct'  
/db\_xref='taxon:32630'  
BASE COUNT 18 a 22 c 29 g 21 t  
ORIGIN

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Best Local Similarity 80.0%; Pred. No. 4.4e+04;  
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 TGCAGATTGCGCAATCTGCA 20  
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Search completed: December 12, 2002, 02:55:49  
Job time.: 321.116 secs

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GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 : Search time 286.304 Seconds  
(without alignments)  
1829.698 Million cell updates/sec

Title: US-09-355-254F-9  
Perfect score: 18  
Sequence: 1 ggaatgacgttcocctgtg 18

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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- 2: gb\_htg.\*
- 3: gb\_in.\*
- 4: gb\_om.\*
- 5: gb\_ov.\*
- 6: gb\_pat.\*
- 7: gb\_ph.\*
- 8: gb\_pl.\*
- 9: gb\_pr.\*
- 10: gb\_ro.\*
- 11: gb\_sts.\*
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- 13: gb\_un.\*
- 14: gb\_vl.\*
- 15: em\_ba.\*
- 16: em\_fun.\*
- 17: em\_hum.\*
- 18: em\_in.\*
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- 22: em\_ov.\*
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- 27: em\_sts.\*
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- 29: em\_vl.\*
- 30: em\_htg\_hum.\*
- 31: em\_htg\_inv.\*
- 32: em\_htg\_other.\*
- 33: em\_htg\_mus.\*
- 34: em\_htg\_pln.\*
- 35: em\_htg\_rtd.\*
- 36: em\_htg\_mam.\*
- 37: em\_htg\_vrt.\*
- 38: em\_sy.\*
- 39: em\_htgo\_hum.\*
- 40: em\_htgo\_mus.\*
- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	18	100.0	18	6	A89788	A89788 Sequence 10
2	18	100.0	18	6	A90875	A90875 Sequence 10
3	18	100.0	18	6	AX455583	AX455583 Sequence 10
C 4	13.8	76.7	38	6	I56082	I56082 Sequence 10
C 5	13.8	76.7	89	6	AX438437	AX438437 Sequence
C 6	13.2	73.3	82	9	HUMMACAJ	L37722 Homo sapien
C 7	13.2	73.3	88	10	MMALDRED04	U89144 Mus musculus
C 8	13	72.2	89	10	MMTCRAC2	X02846 M.musculus
9	12.8	71.1	29	6	AX099976	AX099976 Sequence
10	12.8	71.1	50	6	A16634	A16634 Nucleotide
11	12.8	71.1	50	6	A16645	A16645 Nucleotide
C 12	12.4	68.9	26	6	AX19077	AX19077 Sequence
C 13	12.4	68.9	51	6	AX204048	AX204048 Sequence
C 14	12.4	68.9	67	9	HSWV07A55	U61467 Human myosi
15	12.4	68.9	70	6	AR092212	AR092212 Sequence
16	12.4	68.9	85	6	AX360360	AX360360 Sequence
17	12.2	67.8	30	6	AX469615	AX469615 Sequence
C 18	12.2	67.8	50	6	AX165806	AX165806 Sequence
C 19	12.2	67.8	51	6	AX165000	AX165000 Sequence
C 20	12.2	67.8	89	6	AR045019	AR045019 Sequence
21	12.2	67.8	89	6	I18551	I18551 Sequence 20
22	12.2	67.8	89	6	I34095	I34095 Sequence 20
C 23	12.2	67.8	94	1	ECWITA	X75467 E. coli wit
C 24	12.2	67.8	98	11	MMSTS6	Z36555 M.musculus
C 25	12.2	67.8	98	14	AF264011	AF264011 Hepatitis
C 26	12.2	67.8	100	5	XLRPS4	X64205 X.laavis mr
27	12.2	67.8	100	5	XLRPS11	X64208 X.laavis mr
28	12	66.7	56	6	A43627	AX193249 Sequence
C 29	12	66.7	92	6	AX193249	AX193249 Sequence
C 30	12	66.7	94	6	AX396657	AX396657 Sequence
C 31	11.8	65.6	26	6	I23392	I23392 Sequence 10
C 32	11.8	65.6	28	6	E38332	E38332 Process for
C 33	11.8	65.6	47	6	AX194676	AX194676 Sequence
C 34	11.8	65.6	52	6	E21658	E21658 Spermatogen
C 35	11.8	65.6	74	12	AF405700	AF405700 Synthetic
C 36	11.8	65.6	75	10	MUSIGHCA1	M31028 Mus musculu
C 37	11.8	65.6	77	6	AR081580	AR081580 Sequence
C 38	11.8	65.6	98	6	AX107888	AX107888 Sequence
39	11.6	64.4	25	6	AR089894	AR089894 Sequence
40	11.6	64.4	25	6	AR196929	AR196929 Sequence
C 41	11.6	64.4	25	6	AX116304	AX116304 Sequence
C 42	11.6	64.4	51	6	AX116305	AX116305 Sequence
C 43	11.6	64.4	51	6	AX158271	AX158271 Sequence
44	11.6	64.4	51	6	AX158272	AX158272 Sequence
C 45	11.6	64.4	51	6	AX204448	AX204448 Sequence

ALIGNMENTS

RESULT 1  
A89788  
LOCUS A89788  
DEFINITION Sequence 10 from Patent WO9832462.  
ACCESSION A89788  
VERSION A89788.1 GI:6738302  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unclassified.  
REFERENCE 1 (bases 1 to 18)  
AUTHORS Lipford G.B. and Heeg K.  
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
JOURNAL Patent: WO 9832462-A 10 30-JUL-1998;  
linear DNA 18 bp PAT 22-JAN-2000

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)  
Location/Qualifiers  
1. .18  
/organism="unidentified"  
/db\_xref="taxon:32644"  
3 a 4 c 6 g 5 t

BASE COUNT 3 a 4 c 6 g 5 t  
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Query Match 100.0%; Score 18; DB 6; Length 18;  
Best Local Similarity 100.0%; Pred. No. 23;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18  
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Db 1 GGAATGACGTTCCCTGTG 18

RESULT 2  
A90875  
LOCUS A90875 18 bp DNA linear PAT 22-JAN-2000  
DEFINITION Sequence 10 from Patent EP0855184.  
AUTHORS A90875  
VERSION A90875.1 GI:6739272  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 18)  
AUTHORS Heeg,K.P. and Lipford,G.B.  
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination  
JOURNAL Patent: EP 0855184-A 10 29-JUL-1998;  
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)  
FEATURES  
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/db\_xref="taxon:32644"  
3 a 4 c 6 g 5 t

BASE COUNT 3 a 4 c 6 g 5 t  
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 23;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18  
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Db 1 GGAATGACGTTCCCTGTG 18

RESULT 3  
AX455583  
LOCUS AX455583 18 bp DNA linear PAT 06-JUL-2002  
DEFINITION Sequence 60 from Patent WO222809.  
ACCESSION AX455583  
VERSION AX455583.1 GI:21714651  
KEYWORDS synthetic construct.  
SOURCE synthetic construct.  
ORGANISM artificial sequences.  
REFERENCE 1  
AUTHORS Bauer, S., Lipford, G. and Wagner, H.  
TITLE Process for high throughput screening of cpg-based immuno-agonist/antagonist  
JOURNAL Patent: WO 0222809-A 60 21-MAR-2002;  
Coley Pharmaceutical GmbH (DE)  
FEATURES  
Source 1. .18  
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/note="Synthetic oligonucleotide"  
3 a 4 c 6 g 5 t

BASE COUNT 3 a 4 c 6 g 5 t  
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Best Local Similarity 100.0%; Pred. No. 23;  
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Db 1 GGAATGACGTTCCCTGTG 18

RESULT 4  
I56082/c  
LOCUS I56082 38 bp DNA linear PAT 07-OCT-1997  
DEFINITION Sequence 10 from patent US 5648465.  
ACCESSION I56082  
VERSION I56082.1 GI:2476876  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 38)  
AUTHORS Margolis,R.U., Rauch,U. and Margolis,R.K.  
TITLE Cloning and expression of neurocan, a chondroitin sulfate proteoglycan  
JOURNAL Patent: US 5648465-A 10 15-JUL-1997;  
FEATURES  
Source 1. .38  
/organism="unknown"  
7 a 9 c 5 g 10 t 7 others

BASE COUNT 7 a 9 c 5 g 10 t 7 others  
ORIGIN

Query Match 76.7%; Score 13.8; DB 6; Length 38;  
Best Local Similarity 70.6%; Pred. No. 6.9e+03;  
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QY 1 GGAATGACGTTCCCTGT 17  
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AX438437/c  
LOCUS AX438437 89 bp DNA linear PAT 28-JUN-2002  
DEFINITION Sequence 6852 from Patent WO0229113.  
ACCESSION AX438437  
VERSION AX438437.1 GI:21663245  
KEYWORDS Bacillus clausii.  
SOURCE Bacillus clausii.  
ORGANISM Bacteria; Firmicutes; Bacillales; Bacillaceae; Bacillus.  
REFERENCE 1  
AUTHORS Berka,R. and Clausen,I.G.  
TITLE Methods for monitoring multiple gene expression  
JOURNAL Patent: WO 0229113-A 6852 11-APR-2002;  
Novozymes Biotech, Inc. (US); Novozymes A/S (DK)  
FEATURES  
Source 1. .89  
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/db\_xref="taxon:79880"  
21 a 16 c 29 g 23 t

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ORIGIN

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Matches 15; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 2 GAATGACGTTCCCTGTG 18  
|||||  
Db 88 GAAGACATTCCTGTG 72

RESULT 6  
HUMMACAJ/c  
LOCUS HUMMACAJ 82 bp mRNA linear PRI 11-JAN-1995

```

DEFINITION Homo sapiens (clone 10) macronuclear mRNA.
ACCESSION L37722
VERSION L37722.1 GI:576851
KEYWORDS macronuclear.
SOURCE Homo sapiens cDNA to mRNA.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 82)
AUTHORS Carney,J.P., McKnight,C.E., VanEpps,S. and Kelley,M.R.
TITLE Random amplification of cDNA ends (RRACE) allows for cloning of
multiple novel human cDNA fragments containing CAG repeats
JOURNAL Gene (1994) In press
COMMENT clones were isolated using a CAG oligo. The oligo was made up of
CAG x 8,
so by definition the first 24 bases are CAG x 8 (not included in
the
database entries). The actual number of CAG's is undetermined.
FEATURES
source
1..82
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="10"
/cell_line="Jurkat"
1..82
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20 a 29 c 21 g 10 t 2 others
BASE COUNT 20 a 29 c 21 g 10 t 2 others
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Best Local Similarity 83.3%; Pred. No. 1.5e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
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Db 64 GGCTGACGTTCCCTGTG 47

RESULT 7
MMALDRED04/c
LOCUS Mus musculus aldose reductase gene, exon 4. 88 bp DNA linear ROD 13-APR-1998
DEFINITION Mus musculus aldose reductase gene, exon 4.
ACCESSION U89144
VERSION U89144.1 GI:3046239
KEYWORDS
SEGMENT 4 of 10
SOURCE Mus musculus.
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 88)
AUTHORS McGowan,M.H., Iwata,T. and Carper,D.A.
TITLE Characterization of the mouse aldose reductase gene and promoter in
a lens epithelial cell line
JOURNAL Mol. Vis. 4, 2 (1998)
MEDLINE 98153248
PUBMED 9485485
REMARK http://www.emory.edu/molvis/v4/p2
REFERENCE 2 (bases 1 to 88)
AUTHORS McGowan,M.H., Iwata,T. and Carper,D.A.
TITLE Direct Submission
JOURNAL Submitted (10-FEB-1997) LMOD, National Eye Institute, 9000
Rockville Pike, Bldg6/Rm232, Bethesda, MD 20892, USA
FEATURES
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/organism="Mus musculus"
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21 a 23 c 24 g 20 t
BASE COUNT 21 a 23 c 24 g 20 t
ORIGIN

DEFINITION Homo sapiens (clone 10) macronuclear mRNA.
ACCESSION L37722
VERSION L37722.1 GI:576851
KEYWORDS macronuclear.
SOURCE Homo sapiens cDNA to mRNA.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 82)
AUTHORS Carney,J.P., McKnight,C.E., VanEpps,S. and Kelley,M.R.
TITLE Random amplification of cDNA ends (RRACE) allows for cloning of
multiple novel human cDNA fragments containing CAG repeats
JOURNAL Gene (1994) In press
COMMENT clones were isolated using a CAG oligo. The oligo was made up of
CAG x 8,
so by definition the first 24 bases are CAG x 8 (not included in
the
database entries). The actual number of CAG's is undetermined.
FEATURES
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1. 82
20 a 29 c 21 g 10 t 2 others
BASE COUNT 20 a 29 c 21 g 10 t 2 others
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Query Match 73.3%; Score 13.2; DB 9; Length 82;
Best Local Similarity 83.3%; Pred. No. 1.5e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
II IIIIIIIIIII
Db 64 GGCTGACGTTCCCTGTG 47

RESULT 7
MMALDRED04/c
LOCUS Mus musculus aldose reductase gene, exon 4. 88 bp DNA linear ROD 13-APR-1998
DEFINITION Mus musculus aldose reductase gene, exon 4.
ACCESSION U89144
VERSION U89144.1 GI:3046239
KEYWORDS
SEGMENT 4 of 10
SOURCE Mus musculus.
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 88)
AUTHORS McGowan,M.H., Iwata,T. and Carper,D.A.
TITLE Characterization of the mouse aldose reductase gene and promoter in
a lens epithelial cell line
JOURNAL Mol. Vis. 4, 2 (1998)
MEDLINE 98153248
PUBMED 9485485
REMARK http://www.emory.edu/molvis/v4/p2
REFERENCE 2 (bases 1 to 88)
AUTHORS McGowan,M.H., Iwata,T. and Carper,D.A.
TITLE Direct Submission
JOURNAL Submitted (10-FEB-1997) LMOD, National Eye Institute, 9000
Rockville Pike, Bldg6/Rm232, Bethesda, MD 20892, USA
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21 a 23 c 24 g 20 t
BASE COUNT 21 a 23 c 24 g 20 t
ORIGIN

DEFINITION Homo sapiens (clone 10) macronuclear mRNA.
ACCESSION L37722
VERSION L37722.1 GI:576851
KEYWORDS macronuclear.
SOURCE Homo sapiens cDNA to mRNA.
ORGANISM Homo sapiens
REFERENCE 1 (bases 1 to 82)
AUTHORS Carney,J.P., McKnight,C.E., VanEpps,S. and Kelley,M.R.
TITLE Random amplification of cDNA ends (RRACE) allows for cloning of
multiple novel human cDNA fragments containing CAG repeats
JOURNAL Gene (1994) In press
COMMENT clones were isolated using a CAG oligo. The oligo was made up of
CAG x 8,
so by definition the first 24 bases are CAG x 8 (not included in
the
database entries). The actual number of CAG's is undetermined.
FEATURES
source
1..82
Location/Qualifiers
/organism="Homo sapiens"
/db_xref="taxon:9606"
/clone="10"
/cell_line="Jurkat"
1..82
1. 82
20 a 29 c 21 g 10 t 2 others
BASE COUNT 20 a 29 c 21 g 10 t 2 others
ORIGIN

Query Match 73.3%; Score 13.2; DB 10; Length 88;
Best Local Similarity 83.3%; Pred. No. 1.5e+04;
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18
II IIIIIIIIIII
Db 51 GGTATCACGTTCCCTGAG 34

RESULT 8
MMTCRAC2
LOCUS M.musculus gene for Tcell receptor alpha-chain constant region exon
DEFINITION 2.
ACCESSION X02846
VERSION X02846.1 GI:54472
KEYWORDS constant region; T-cell receptor.
SOURCE Mus musculus.
ORGANISM Mus musculus
REFERENCE 1 (bases 1 to 89)
AUTHORS Hayday,A.C., Diamond,D.J., Tanigawa,G., Heilig,J.S., Folsom,V.,
Saito,H. and Tonegawa,S.
TITLE Unusual organization and diversity of T-cell receptor alpha-chain
genes
JOURNAL Nature 316 (6031), 828-832 (1985)
MEDLINE 85296331
PUBMED 2993907
COMMENT The coding sequence is identical to that of alpha-chain mRNA
isolated from cytotoxic T-cells (clone PHDS58, see X01134). The
first base of the first codon of PHDS58 constant region is
identical to pos. 91 in X02843.
FEATURES
source
1..89
/organism="Mus musculus"
/db_xref="taxon:10090"
/cell_type="lymphocyte T"
<1..21
/note="intron"
22..66
/product="constant region of Tcell alpha chain"
/note="exon 2"
67..>89
/note="intron"
23 a 20 c 21 g 25 t
BASE COUNT 23 a 20 c 21 g 25 t
ORIGIN

Query Match 72.2%; Score 13; DB 10; Length 89;
Best Local Similarity 100.0%; Pred. No. 2e+04;
Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 GACGTTCCCTGTG 18
IIIIIIIIIIII
Db 21 GACGTTCCCTGTG 33

RESULT 9
AX099976
LOCUS Sequence 5 from Patent WO0120007.
DEFINITION AX099976
ACCESSION AX099976
VERSION AX099976.1 GI:13538986
KEYWORDS synthetic construct.
SOURCE synthetic construct
ORGANISM artificial sequences.
REFERENCE 1 (bases 1 to 29)
AUTHORS Sillaots,S., Martinez-Perez,A., Tsang,A. and Storms,R.
TITLE A multifunctional system for the efficient manipulation of protein
expression in filamentous fungi and method using same
JOURNAL Patent: WO 0120007-A 5 22-MAR-2001;
Concordia University (CA)

```

Query Match	71.1%	Score 12.8;	DB 6;	Length 50;
Best Local Similarity	87.5%	Pred. No. 2.7e+04;		
Matches 14; Conservative	0;	Mismatches 2;	Indels 0;	Gaps 0;

  

QY	1	GGAATGACGTTCCCTG 16		
DB	5	GGAATGACGATCCCTG 20		

  

RESULT 12				
AX419077/c				
LOCUS	AX419077	26 bp	DNA	linear
DEFINITION	Sequence 96 from Patent WO0212471.			
ACCESSION	AX419077			
VERSION	AX419077.1	GI:21523851		
KEYWORDS				
SOURCE	synthetic construct.			
ORGANISM	artificial construct			
REFERENCE	1			
AUTHORS	Acton, S., Robison, K.E. and Hsieh, F.Y.			
TITLE	Angiotensin converting enzyme homolog and uses therefor			
JOURNAL	Patent: WO 0212471-A 96 14-FEB-2002;			
	Millennium Pharmaceuticals, Inc. (US)			
FEATURES	Location/Qualifiers			
source	1..26			
	/organism="synthetic construct"			
	/db_xref="taxon:32630"			
	/note="motifs"			
BASE COUNT	7 a	3 c	9 g	7 t
ORIGIN				

  

Query Match	68.9%	Score 12.4;	DB 6;	Length 26;
Best Local Similarity	92.9%	Pred. No. 4.8e+04;		
Matches 13; Conservative	0;	Mismatches 1;	Indels 0;	Gaps 0;

  

QY	2	GAATGACGTTCCCT 15		
DB	16	GAATGACTTCCCT 3		

  

RESULT 13				
AX204048/c				
LOCUS	AX204048	51 bp	DNA	linear
DEFINITION	Sequence 154 from Patent WO0148245.			
ACCESSION	AX204048			
VERSION	AX204048.1	GI:15393526		
KEYWORDS	human.			
SOURCE	Homo sapiens			
ORGANISM	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.			
REFERENCE	1	(bases 1 to 51)		
AUTHORS	Shinkets, R.A. and Leach, M.			
TITLE	Nucleic acids containing single nucleotide polymorphisms and methods of use thereof			
JOURNAL	Patent: WO 0148245-A 154 05-JUL-2001;			
	Curagen Corporation (US)			
FEATURES	Location/Qualifiers			
source	1..51			
	/organism="Homo sapiens"			
	/db_xref="taxon:9606"			
variation	26			
	/note="single nucleotide polymorphism"			
	Accession number cg44000740"			
BASE COUNT	16 a	15 c	13 g	7 t
ORIGIN				

  

Query Match	68.9%	Score 12.4;	DB 6;	Length 51;
Best Local Similarity	92.9%	Pred. No. 4.6e+04;		
Matches 13; Conservative	0;	Mismatches 1;	Indels 0;	Gaps 0;

QY 5 TGACGTTCCCTGTG 18  
| | | | | | | | | |  
Db 39 TGATGTTCCCTGTG 26

RESULT 14  
HSMY07A55 HSMY07A55 67 bp DNA linear PRI 19-MAR-1997  
LOCUS Human myosin VIIa (MYO7A) gene, 5' exon 38.  
DEFINITION U61467  
ACCESSION U61467  
VERSION U61467.1 GI:1894878

KEYWORDS 55 of 77  
SEGMENT Homo sapiens.  
SOURCE Homo sapiens  
ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE 1 (bases 1 to 67)  
AUTHORS Kelley.P.M., Weston.M.D., Chen.Z.-Y., Orten.D.J., Hasson.T.,  
Overbeck.L.D., Pinnti.J., Talmadge.C.B., Ing.P., Mooseker.M.S.,  
Corey.D., Sumegi.J. and Kimberling.W.J.

TITLE The genomic structure of the gene defective in Usher syndrome type  
Ib (MYO7A)

JOURNAL Genomics 40 (1), 73-79 (1997)

MEDLINE 97224487

PUBMED 9070921

REFERENCE 2 (bases 1 to 67)

AUTHORS Kelley.P.M.

TITLE Direct Submission

JOURNAL Submitted (21-JUN-1996) Gene Marker Lab, Boys Town National  
Research Hospital, 555 North 30th Street, Omaha, NE 68131, USA

FEATURES Location/Qualifiers

source 1..67

exon /organism="Homo sapiens"

42..>67 /db\_xref="taxon:9606"

/gene="MYO7A"

/number=38

BASE COUNT 11 a 24 c 21 g 11 t

ORIGIN

Query Match 68.9%; Score 12.4; DB 9; Length 67;

Best Local Similarity 92.9%; Pred. No. 4.5e+04;

Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 5 TGACGTTCCCTGTG 18  
| | | | | | | | | |  
Db 19 TGACGTTCCCTGTG 32

RESULT 15  
AR092212 AR092212 70 bp DNA linear PAT 08-SEP-2000  
LOCUS Sequence 150 from patent US 598142.  
DEFINITION AR092212  
ACCESSION AR092212  
VERSION AR092212.1 GI:10018966

KEYWORDS Unknown.  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 70)  
AUTHORS Gold,L., Eaton,B., Smith,D., Wecker,M. and Jensen,K.  
TITLE Systematic evolution of ligands by exponential enrichment:  
chemi-SELEX  
JOURNAL Patent: US 598142-A 150 07-DEC-1999;

FEATURES Location/Qualifiers

source 1..70

BASE COUNT 16 a 17 c 25 g 12 t

ORIGIN /organism="unknown"

Query Match 68.9%; Score 12.4; DB 6; Length 70;

Best Local Similarity 92.9%; Pred. No. 4.5e+04;

Matches 13; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AATGACGTTCCCTG 16

| | | | | | | | | |

Db 34 AATGACGTACCTG 47

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Job time : 295.304 secs

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score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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(without alignments)  
1829.698 Million cell updates/sec

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Sequence: 1 gtccattcccgtaaatctt 20

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

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Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Result No.	Score	Query Match	Length	DB ID	Description
1	20	100.0	20	6	A89795 Sequence 17
2	20	100.0	20	6	A90882 Sequence 17
3	20	100.0	20	6	AX455587 Sequence
4	17.4	87.0	22	6	AX040434 Sequence
5	17	85.0	19	6	I81948 Sequence 46
6	15.4	77.0	17	6	I87792 Sequence 20
7	15.4	77.0	19	6	I81949 Sequence 47
8	15.4	77.0	22	6	AX040435 Sequence
9	15.4	77.0	25	6	AR182575 Sequence
10	15.4	77.0	25	6	AX472568 Sequence
11	15.4	77.0	25	6	AX476844 Sequence
12	15.4	77.0	25	6	AX476862 Sequence
13	15.4	77.0	28	6	I39727 Sequence 14
14	15.4	77.0	28	6	I55844 Sequence 14
15	15.2	76.0	20	6	AR103469 Sequence
16	15.2	76.0	20	6	AR176473 Sequence
17	14.4	72.0	18	6	AR153598 Sequence
18	14.2	71.0	26	6	AR160395 Sequence
19	14.2	71.0	26	6	AX482621 Sequence
20	14.2	71.0	72	6	I95007 Sequence 24
21	14.2	71.0	76	6	I95006 Sequence 23
22	14	70.0	19	6	AX386646 Sequence
23	13.8	69.0	25	6	AX192408 Sequence
24	13.8	69.0	28	6	I39716 Sequence 3
25	13.8	69.0	28	6	I55833 Sequence 3
26	13.6	68.0	35	6	AR061506 Sequence
27	13.6	68.0	35	6	AR108405 Sequence
28	13.6	68.0	35	6	I16362 Sequence 18
29	13.6	68.0	35	6	I66848 Sequence 18
30	13.6	68.0	35	6	I84942 Sequence 18
31	13.6	68.0	94	11	AL773118 Arabidops
32	13.2	66.0	66	10	MUSMUP
33	13.2	66.0	83	3	DME428807
34	13	65.0	13	6	AX026536 Sequence
35	12.8	64.0	52	6	AR103470 Sequence
36	12.8	64.0	52	6	AR176474 Sequence
37	12.6	63.0	22	6	AR062163 Sequence
38	12.6	63.0	30	6	AR004706 Sequence
39	12.6	63.0	30	6	AR008192 Sequence
40	12.6	63.0	30	6	AR136975 Sequence
41	12.6	63.0	30	6	I76976 Sequence 36
42	12.6	63.0	30	6	I80971 Sequence 36
43	12.6	63.0	30	6	I81067 Sequence 36
44	12.6	63.0	46	6	AR126436 Sequence
45	12.6	63.0	50	6	AR135509 Sequence

ALIGNMENTS

RESULT 1  
A89795  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL

A89795  
Sequence 17 from Patent WO9832462.  
A89795  
A89795.1 GI:6738309  
unidentified.  
unidentified.  
1 (bases 1 to 20)  
Lipford,G.B. and Heeg,K.  
PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
Patent: WO 9832462-A 17 30-JUL-1998

20 bp  
DNA  
linear  
PAT 22-JAN-2000

```

FEATURES
Source
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
Location/Qualifiers
1..20
/organism="unidentified"
/db_xref="taxon:32644"
BASE COUNT      4 a      6 c      2 g      8 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCCATTTCCTCGTAATCTT 20
    |||
Db 1 GTCCATTTCCTCGTAATCTT 20
    |||

RESULT 2
AX455587
LOCUS
DEFINITION
Sequence 64 from Patent WO222809.
ACCESSION
AX455587
VERSION
AX455587.1 GI:21714655
KEYWORDS
synthetic construct.
SOURCE
artificial sequences.
ORGANISM
Bauer, S., Lipford, G. and Wagner, H.
REFERENCE
1 (bases 1 to 20)
AUTHORS
Heeg, K.P. and Lipford, G.B.
TITLE
Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL
Patent: EP 0855184-A 17 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES
Location/Qualifiers
1..20
/organism="unidentified"
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BASE COUNT      4 a      6 c      2 g      8 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCCATTTCCTCGTAATCTT 20
    |||
Db 1 GTCCATTTCCTCGTAATCTT 20
    |||

RESULT 3
AX455587
LOCUS
DEFINITION
Sequence 64 from Patent WO222809.
ACCESSION
AX455587
VERSION
AX455587.1 GI:21714655
KEYWORDS
synthetic construct.
SOURCE
artificial sequences.
ORGANISM
Bauer, S., Lipford, G. and Wagner, H.
REFERENCE
1 (bases 1 to 20)
AUTHORS
Heeg, K.P. and Lipford, G.B.
TITLE
Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL
Patent: EP 0855184-A 17 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES
Location/Qualifiers
1..20
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BASE COUNT      4 a      6 c      2 g      8 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 6;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTCCATTTCCTCGTAATCTT 20
    |||
Db 1 GTCCATTTCCTCGTAATCTT 20
    |||

RESULT 4
AX404034
LOCUS
DEFINITION
Sequence 9 from Patent WO063357.
ACCESSION
AX404034
VERSION
AX404034.1 GI:11230241
KEYWORDS
synthetic construct.
SOURCE
artificial sequences.
ORGANISM
Flier, J.S. and Bjorbaek, C.
REFERENCE
1 (bases 1 to 22)
AUTHORS
Methods and compositions for modulating ciliary neurotrophic facto
TITLE
r activity
JOURNAL
Patent: WO 0063357-A 9 26-OCT-2000;
Beth Israel Deaconess Medical Center (US)
FEATURES
Location/Qualifiers
1..22
/organism="synthetic construct"
/db_xref="taxon:32630"
/note="primer-bind"
BASE COUNT      5 a      8 c      2 g      7 t
ORIGIN

Query Match      87.0%; Score 17.4; DB 6; Length 22;
Best Local Similarity 94.7%; Pred. No. 1.5e+02;
Matches 18; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCATTTCCTCGTAATCTT 20
    |||
Db 4 TCATTTCCTCGTAATCAT 22
    |||

RESULT 5
I81948
LOCUS
DEFINITION
Sequence 46 from patent US 5712094.
ACCESSION
I81948
VERSION
I81948.1 GI:3210245
KEYWORDS
Unknown.
SOURCE
Unclassified.
ORGANISM
Seidel, H. Martin., Lamb, I. Peter., and Chan, S.-S. Tian.
REFERENCE
1 (bases 1 to 19)
AUTHORS
Methods for detecting modulators of cytokine action
TITLE
Patent: US 5712094-A 46 27-JAN-1998;
JOURNAL
Location/Qualifiers
1..19
FEATURES
/organism="unknown"
BASE COUNT      5 a      6 c      2 g      6 t
ORIGIN

Query Match      85.0%; Score 17; DB 6; Length 19;
Best Local Similarity 100.0%; Pred. No. 2.3e+02;
Matches 17; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 2 TCATTTCCTCGTAATCTT 18
    |||
Db 3 TCATTTCCTCGTAATCTT 19
    |||

RESULT 6
I87792
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FEATURES	source	Location/Qualifiers
BASE COUNT.	6 a 4 c 6 g 6 t	
ORIGIN		
Query Match	77.0%; Score 15.4; DB 6; Length 22;	
Best Local Similarity	94.1%; Pred. No. 1.7e+03;	
Matches	16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
QY	4 CATTTCCTCGTAAATCTT 20	
DB		
LOCUS	AR182575	25 bp DNA linear PAT 20-APR-2002
DEFINITION	Sequence 23 from patent US 6338949.	
ACCESSION	AR182575	
VERSION	AR182575.1 GI:20225782	
KEYWORDS	Unknown.	
SOURCE	Unknown.	
ORGANISM	Unknown.	
REFERENCE	1 (bases 1 to 25)	
AUTHORS	Darnell,J.E. Jr., Schindler,C.W., Fu,X.-Y., Wen,Z. and Zhong,Z.	
TITLE	Nucleic acids encoding receptor recognition factor stat4 and methods of use thereof	
JOURNAL	Patent: US 6338949-A 23 15-JAN-2002;	
FEATURES	Location/Qualifiers	
source	1..25	
BASE COUNT	6 a 7 c 5 g 7 t	
ORIGIN	/organism="unknown"	
Query Match	77.0%; Score 15.4; DB 6; Length 25;	
Best Local Similarity	94.1%; Pred. No. 1.7e+03;	
Matches	16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;	
QY	4 CATTTCCTCGTAAATCTT 20	
DB		
LOCUS	AX472568	25 bp DNA linear PAT 09-AUG-2002
DEFINITION	Sequence 63 from Patent WO02052039.	
ACCESSION	AX472568	
VERSION	AX472568.1 GI:22207472	
KEYWORDS	synthetic construct.	
SOURCE	synthetic construct	
ORGANISM	artificial sequences.	
REFERENCE	1	
AUTHORS	Blais,Y., Rousseau,P., Leblanc,B. and Camato,R.N.	
TITLE	Methods for selecting and producing selective pharmaceutical compounds and compositions using an established genetically altered cell-based library responsive to transcription factors; genetic constructs and library therefor	
JOURNAL	Patent: WO 02052039-A 63 04-JUL-2002;	
FEATURES	Geneka Biotechnology Inc. (CA)	
source	Location/Qualifiers	
BASE COUNT	6 a 7 c 5 g 7 t	
ORIGIN	/organism="synthetic construct"	
	/db_xref="taxon:32630"	
	/note="Oligonucleotide"	

Query Match 77.0%; Score 15.4; DB 6; Length 25;  
Best Local Similarity 94.1%; Pred. No. 1.7e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CATTTCCTCGTAATCTT 20  
|||||  
Db 6 CATTTCCTCGTAATCTG 22

RESULT 11  
AX476844  
LOCUS AX476844 25 bp DNA linear PAT 12-AUG-2002  
DEFINITION Sequence 21 from Patent WO02052037.  
ACCESSION AX476844  
VERSION AX476844.1 GI:22216120  
KEYWORDS synthetic construct.  
SOURCE synthetic construct  
ORGANISM artificial sequences.  
REFERENCE 1  
AUTHORS Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.  
TITLE Method for screening and/or identifying factors that bind to nucleic acids  
JOURNAL Patent: WO 02052037-A 21 04-JUL-2002;  
Geneka Biotechnology Inc. (CA)  
FEATURES Location/Qualifiers  
source  
1..25  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="NABE-probes"

BASE COUNT 6 a 7 c 5 g 7 t  
ORIGIN

Query Match 77.0%; Score 15.4; DB 6; Length 25;  
Best Local Similarity 94.1%; Pred. No. 1.7e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CATTTCCTCGTAATCTT 20  
|||||  
Db 6 CATTTCCTCGTAATCTG 22

RESULT 12  
AX476862  
LOCUS AX476862 25 bp DNA linear PAT 12-AUG-2002  
DEFINITION Sequence 39 from Patent WO02052037.  
ACCESSION AX476862  
VERSION AX476862.1 GI:22216138  
KEYWORDS synthetic construct.  
SOURCE synthetic construct  
ORGANISM artificial sequences.  
REFERENCE 1  
AUTHORS Larose,A.M., Rousseau,P., Leblanc,B. and Camato,R.  
TITLE Method for screening and/or identifying factors that bind to nucleic acids  
JOURNAL Patent: WO 02052037-A 39 04-JUL-2002;  
Geneka Biotechnology Inc. (CA)  
FEATURES Location/Qualifiers  
source  
1..25  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="Double stranded NABE"

BASE COUNT 6 a 7 c 5 g 7 t  
ORIGIN

Query Match 77.0%; Score 15.4; DB 6; Length 25;  
Best Local Similarity 94.1%; Pred. No. 1.7e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 4 CATTTCCTCGTAATCTT 20  
|||||  
Db 6 CATTTCCTCGTAATCTG 22

RESULT 13  
I39727  
LOCUS I39727 28 bp DNA linear PAT 13-MAY-1997  
DEFINITION Sequence 14 from patent US 5616489.  
ACCESSION I39727  
VERSION I39727.1 GI:2084207  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 28)  
AUTHORS Levy,D.E.  
TITLE DNA sequence which binds transcriptional regulatory proteins activated in response to various cytokines and uses thereof  
JOURNAL Patent: US 5616489-A 14 01-APR-1997;  
FEATURES Location/Qualifiers  
source  
1..28  
/organism="unknown"

BASE COUNT 7 a 9 c 3 g 9 t  
ORIGIN

Query Match 77.0%; Score 15.4; DB 6; Length 28;  
Best Local Similarity 94.1%; Pred. No. 1.7e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCCATTTCCTCGTAATC 18  
|||||  
Db 8 TTCATTTCCTCGTAATC 24

RESULT 14  
I55844  
LOCUS I55844 28 bp DNA linear PAT 07-OCT-1997  
DEFINITION Sequence 14 from patent US 5648217.  
ACCESSION I55844  
VERSION I55844.1 GI:2476638  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 28)  
AUTHORS Levy,D.E.  
TITLE DNA sequence which binds transcriptional regulatory proteins activated in response to various cytokines and uses thereof  
JOURNAL Patent: US 5648217-A 14 15-JUL-1997;  
FEATURES Location/Qualifiers  
source  
1..28  
/organism="unknown"

BASE COUNT 7 a 9 c 3 g 9 t  
ORIGIN

Query Match 77.0%; Score 15.4; DB 6; Length 28;  
Best Local Similarity 94.1%; Pred. No. 1.7e+03;  
Matches 16; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 2 TCCATTTCCTCGTAATC 18  
|||||  
Db 8 TTCATTTCCTCGTAATC 24

RESULT 15  
AR103469  
LOCUS AR103469 20 bp DNA linear PAT 14-FEB-2001  
DEFINITION Sequence 5 from patent US 6087478.  
ACCESSION AR103469  
VERSION AR103469.1 GI:12815057  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.

REFERENCE 1 (bases 1 to 20)

AUTHORS Vinkemeier, U., Moarefi, I., Darnell, J.E. Jr. and Kuriyan, J.  
 TITLE Crystal of the N-terminal domain of a STAT protein and methods of  
 use thereof  
 JOURNAL Patent: US 6087478-A 5 11-JUL-2000;  
 FEATURES Location/Qualifiers  
 source 1..20  
 BASE COUNT 5 a 6 c 3 g 6 t  
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 Best Local Similarity 85.0%; Pred. No. 2.1e+03;  
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 Db 1 GCCGATTTCCTGTAATCAT 20  
 Search completed: December 12, 2002, 02:55:59  
 Job time : 323.116 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model  
Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds  
(without alignments)  
1829.698 Million cell updates/sec

Title: US-09-355-254F-17  
Perfect score: 20  
Sequence: 1 tatgcataattctgtaagtg 20  
Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues  
Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

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2	20	100.0	20	6	A90883	A90883 Sequence 18
3	20	100.0	20	6	AX455591	AX455591 Sequence
4	19	95.0	24	6	AX406780	AX406780 Sequence
5	16	80.0	19	6	I79489	I79489 Sequence 75
6	16	80.0	19	6	I81942	I81942 Sequence 40
7	15	75.0	17	6	I87793	I87793 Sequence 21
C 8	15	75.0	19	6	I79490	I79490 Sequence 76
C 9	15	75.0	19	6	I81943	I81943 Sequence 41
C 10	14.8	74.0	47	6	AR032425	AR032425 Sequence
C 11	14.8	74.0	47	6	AR209089	AR209089 Sequence
C 12	14.8	74.0	47	6	I29165	I29165 Sequence 37
C 13	14.8	74.0	47	6	I90839	I90839 Sequence 37
C 14	14.8	74.0	50	6	AR032824	AR032824 Sequence
C 15	14.8	74.0	50	6	AR032825	AR032825 Sequence
C 16	14.8	74.0	50	6	AR209488	AR209488 Sequence
C 17	14.8	74.0	50	6	AR209489	AR209489 Sequence
C 18	14.8	74.0	50	6	I29564	I29564 Sequence 43
C 19	14.8	74.0	50	6	I29565	I29565 Sequence 43
C 20	14.8	74.0	50	6	I91238	I91238 Sequence 43
C 21	14.8	74.0	50	6	I91239	I91239 Sequence 43
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23	14.4	72.0	19	6	I79439	I79439 Sequence 25
24	14.4	72.0	19	6	I79447	I79447 Sequence 33
25	14.4	72.0	19	6	I79463	I79463 Sequence 49
26	14.4	72.0	19	6	I79465	I79465 Sequence 51
27	14.4	72.0	19	6	I79469	I79469 Sequence 55
28	14.4	72.0	19	6	I81944	I81944 Sequence 42
29	14.4	72.0	32	6	E09298	E09298 DNA linker
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C 33	13.4	67.0	19	6	I79448	I79448 Sequence 34
C 34	13.4	67.0	19	6	I79464	I79464 Sequence 50
C 35	13.4	67.0	19	6	I79466	I79466 Sequence 52
C 36	13.4	67.0	19	6	I79470	I79470 Sequence 56
C 37	13.4	67.0	19	6	I81945	I81945 Sequence 43
C 38	13.4	67.0	24	6	AX444661	AX444661 Sequence
C 39	13.4	67.0	25	6	AX116799	AX116799 Sequence
40	13.4	67.0	51	6	AX163145	AX163145 Sequence
41	13.4	67.0	60	6	ARI77644	ARI77644 Sequence
42	13.4	67.0	93	9	AF281956	AF281956 Homo sapi
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ALIGNMENTS

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ACCESSION A89796  
VERSION A89796.1 GI:6738310  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Lipford, G. B. and Heeg, K.  
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
JOURNAL Patent: WO 9832462-A 18 30-JUL-1998;  
linear PAT 22-JAN-2000

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RESULT 2
AX0883
LOCUS
DEFINITION      A90883
Sequence 18 from Patent EP0855184.
ACCESSION      A90883
VERSION
A90883.1 GI:6739312
KEYWORDS
SOURCE
ORGANISM
unidentified.
unidentified.
unclassified.
REFERENCE
1 (bases 1 to 20)
AUTHORS
Heeg,K.P. and Lipford,G.B.
TITLE
Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL
Patent: EP 0855184-A 18 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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RESULT 3
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LOCUS
DEFINITION      AX455591
Sequence 68 from Patent WO0222809.
ACCESSION      AX455591
VERSION
AX455591.1 GI:21714659
KEYWORDS
SOURCE
synthetic construct.
artificial sequences.
ORGANISM
1
REFERENCE
Bauer,S., Lipford,G. and Wagner,H.
AUTHORS
Process for high throughput screening of cpg-based
immuno-agonist/antagonist
TITLE
Patent: WO 0222809-A 68 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)
JOURNAL
Coley Pharmaceutical GmbH (DE)
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/Note="Synthetic oligonucleotide"
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Db 1 TATGCATATTCCTGTAAGTG 20

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Sequence 36 from Patent WO0229044.
ACCESSION      AX406780
VERSION
AX406780.1 GI:21439705
KEYWORDS
SOURCE
synthetic construct.
synthetic construct
artificial sequences.
ORGANISM
1
REFERENCE
Hecker,M. and Wagner,A.H.
AUTHORS
Modulation of the transcription of pro-inflammatory gene products
TITLE
Patent: WO 0229044-A 36 11-APR-2002;
JOURNAL
Location/Qualifiers
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Db 6 TATGCATATTCCTGTAAGT 24

RESULT 5
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LOCUS
DEFINITION      I79489
Sequence 75 from patent US 5707803.
ACCESSION      I79489
VERSION
I79489.1 GI:3207779
KEYWORDS
SOURCE
Unknown.
ORGANISM
Unclassified.
REFERENCE
1 (bases 1 to 19)
AUTHORS
Lamb,I.Peter. and Seidel,H.Martin.
TITLE
DNA regulatory elements responsive to cytokines and methods for
their use
JOURNAL
Patent: US 5707803-A 75 13-JAN-1998;
FEATURES
Source
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Location/Qualifiers
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Db 4 CATATTTCCTGTAAGTG 19

RESULT 6
I81942
LOCUS

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ACCESSION I81942  
VERSION I81942.1 GI:3210239  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE 1 (bases 1 to 19)  
AUTHORS Seidel, H. Martin., Lamb, I. Peter. and Chan, S.-S. Tian.  
TITLE Methods for detecting modulators of cytokine action  
JOURNAL Patent: US 5712094-A 40 27-JAN-1998;  
FEATURES Location/Qualifiers  
source 1..19  
BASE COUNT 5 a 3 c 4 g 7 t  
ORIGIN

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Best Local Similarity 100.0%; Pred. No. 3.2e+03;  
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DEFINITION Sequence 21 from patent US 5716622.  
ACCESSION I87793  
VERSION I87793.1 GI:3407733  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE 1 (bases 1 to 17)  
AUTHORS Darnell, J.E. Jr., Wen, Z., Horvath, C.M. and Zhong, Z.  
TITLE Functionally active regions of signal transducer and activators of transcription  
JOURNAL Patent: US 5716622-A 21 10-FEB-1998;  
FEATURES Location/Qualifiers  
source 1..17  
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RESULT 8  
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ACCESSION I79490  
VERSION I79490.1 GI:3207780  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE 1 (bases 1 to 19)  
AUTHORS Lamb, I. Peter. and Seidel, H. Martin.  
TITLE DNA regulatory elements responsive to cytokines and methods for their use  
JOURNAL Patent: US 5707803-A 76 13-JAN-1998;  
FEATURES Location/Qualifiers  
source 1..19

Query Match 75.0%; Score 15; DB 6; Length 19;  
Best Local Similarity 100.0%; Pred. No. 9.9e+03;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20  
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Db 19 ATATTCCTGTAAGTG 5

RESULT 9  
I81943/c  
LOCUS  
DEFINITION Sequence 41 from patent US 5712094.  
ACCESSION I81943  
VERSION I81943.1 GI:3210240  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE 1 (bases 1 to 19)  
AUTHORS Seidel, H. Martin., Lamb, I. Peter. and Chan, S.-S. Tian.  
TITLE Methods for detecting modulators of cytokine action  
JOURNAL Patent: US 5712094-A 41 27-JAN-1998;  
FEATURES Location/Qualifiers  
source 1..19  
BASE COUNT 7 a 4 c 3 g 5 t  
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Db 19 ATATTCCTGTAAGTG 5

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ACCESSION AR032425  
VERSION AR032425.1 GI:5948030  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE 1 (bases 1 to 47)  
AUTHORS Edwards, C.A., Cantor, C.R., Andrews, B.M., Turin, L.M. and Fry, K.E.  
TITLE Method of determining DNA sequence preference of a DNA-binding molecule  
JOURNAL Patent: US 5869241-A 37 09-FEB-1999;  
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BASE COUNT 17 a 6 c 10 g 14 t  
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DEFINITION Sequence 37 from patent US 6384208.  
ACCESSION AR209089  
VERSION AR209089.1 GI:21510414  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE  
1 (bases 1 to 47)  
Unclassified.  
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.  
TITLE Sequence directed DNA binding molecules compositions and methods  
JOURNAL Patent: US 6384208-A 37 07-MAY-2002;  
FEATURES  
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BASE COUNT 17 a 6 c 10 g 14 t  
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Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

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Db 28 TATTTATATTCCTGTAAG 11

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DEFINITION Sequence 37 from patent US 5578444.  
ACCESSION I29165  
VERSION I29165.1 GI:1819956  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE  
1 (bases 1 to 47)  
Unclassified.  
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.  
TITLE Sequence-directed DNA-binding molecules compositions and methods  
JOURNAL Patent: US 5578444-A 37 26-NOV-1996;  
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BASE COUNT 17 a 6 c 10 g 14 t  
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Db 28 TATTTATATTCCTGTAAG 11

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ACCESSION I90839  
VERSION I90839.1 GI:3935309  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE  
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Unclassified.  
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M. and Turin,L.M.  
TITLE Screening assay for the detection of DNA-binding molecules  
JOURNAL Patent: US 5726014-A 37 10-MAR-1998;  
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DEFINITION Sequence 436 from patent US 5869241.  
ACCESSION AR032824  
VERSION AR032824.1 GI:5948429  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE  
1 (bases 1 to 50)  
Unclassified.  
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.  
TITLE Method of determining DNA sequence preference of a DNA-binding molecule.  
JOURNAL Patent: US 5869241-A 436 09-FEB-1999;  
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Location/Qualifiers  
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BASE COUNT 18 a 6 c 10 g 16 t  
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DEFINITION Sequence 437 from patent US 5869241.  
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VERSION AR032825.1 GI:5948430  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE  
1 (bases 1 to 50)  
Unclassified.  
AUTHORS Edwards,C.A., Cantor,C.R., Andrews,B.M., Turin,L.M. and Fry,K.E.  
TITLE Method of determining DNA sequence preference of a DNA-binding molecule.  
JOURNAL Patent: US 5869241-A 437 09-FEB-1999;  
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Job time : 321.116 secs

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LOCUS A90888 20 bp DNA linear PAT 22-JAN-2000  
DEFINITION Sequence 23 from Patent EP0855184.  
ACCESSION A90888  
VERSION A90888.1 GI:6739338  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unclassified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Heeg,K.P. and Lipford,G.B.  
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination  
JOURNAL Patent: EP 0855184-A 23 29-JUL-1998;  
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)  
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DEFINITION Sequence 22 from Patent WO0222809.  
ACCESSION AX455545  
VERSION AX455545.1 GI:21714613  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.  
REFERENCE 1  
AUTHORS Bauer,S., Lipford,G. and Wagner,H.  
TITLE Process for high throughput screening of cpqg-based immuno-agonist/antagonist  
JOURNAL Patent: WO 0222809-A 22 21-MAR-2002;  
Coley Pharmaceutical GmbH (DE)  
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Db 1 AAGCGAAATGAATTGACT 20

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DEFINITION Sequence 19 from Patent WO0229044.  
ACCESSION AX406763  
VERSION AX406763.1 GI:21439688  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.  
REFERENCE 1  
AUTHORS Hecker,M. and Wagner,A.H.  
TITLE Modulation of the transcription of pro-inflammatory gene products  
JOURNAL Patent: WO 0229044-A 19 11-APR-2002;  
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Db 3 AAGCGAAATGAATTGACT 22

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LOCUS AX406764 22 bp DNA linear PAT 15-JUN-2002  
DEFINITION Sequence 20 from Patent WO0229044.  
ACCESSION AX406764  
VERSION AX406764.1 GI:21439689  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.  
REFERENCE 1  
AUTHORS Hecker,M. and Wagner,A.H.  
TITLE Modulation of the transcription of pro-inflammatory gene products  
JOURNAL Patent: WO 0229044-A 20 11-APR-2002;  
FEATURES Location/Qualifiers  
source 1..22  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="oligonucleotide"  
BASE COUNT 4 a 6 c 2 g 10 t  
ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 22;  
Best Local Similarity 100.0%; Pred. No. 1.2e+02;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCGAAATGAATTGACT 20  
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Db 20 AAGCGAAATGAATTGACT 1

RESULT 6  
AX482985/c

Query Match 100.0%; Score 20; DB 6; Length 20;  
Best Local Similarity 100.0%; Pred. No. 1.2e+02;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCGAAATGAATTGACT 20  
|||||  
Db 1 AAGCGAAATGAATTGACT 20

RESULT 4  
AX406763  
LOCUS AX406763 22 bp DNA linear PAT 14-JUN-2002  
DEFINITION Sequence 19 from Patent WO0229044.  
ACCESSION AX406763  
VERSION AX406763.1 GI:21439688  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.  
REFERENCE 1  
AUTHORS Hecker,M. and Wagner,A.H.  
TITLE Modulation of the transcription of pro-inflammatory gene products  
JOURNAL Patent: WO 0229044-A 19 11-APR-2002;  
FEATURES Location/Qualifiers  
source 1..22  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="oligonucleotide"  
BASE COUNT 10 a 2 c 6 g 4 t  
ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 22;  
Best Local Similarity 100.0%; Pred. No. 1.2e+02;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCGAAATGAATTGACT 20  
|||||  
Db 3 AAGCGAAATGAATTGACT 22

RESULT 5  
AX406764/c  
LOCUS AX406764 22 bp DNA linear PAT 15-JUN-2002  
DEFINITION Sequence 20 from Patent WO0229044.  
ACCESSION AX406764  
VERSION AX406764.1 GI:21439689  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.  
REFERENCE 1  
AUTHORS Hecker,M. and Wagner,A.H.  
TITLE Modulation of the transcription of pro-inflammatory gene products  
JOURNAL Patent: WO 0229044-A 20 11-APR-2002;  
FEATURES Location/Qualifiers  
source 1..22  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
/note="oligonucleotide"  
BASE COUNT 4 a 6 c 2 g 10 t  
ORIGIN

Query Match 100.0%; Score 20; DB 6; Length 22;  
Best Local Similarity 100.0%; Pred. No. 1.2e+02;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 AAGCGAAATGAATTGACT 20  
|||||  
Db 20 AAGCGAAATGAATTGACT 1

RESULT 6  
AX482985/c

20  
/note="single nucleotide polymorphism  
Accession number cg43119818"

Database	Accession	Length	Source
Db	22 AACCAAAATCAATTCAGT	3	LOCUS
	22 AACCAAAATCAATTCAGT	3	HUMADPRT11
		89 bp	DNA
		1 linear	PRI 30-OCT-1994



DEFINITION Human NAD+ ADP-ribosyltransferase (ADPRT) gene, exon 11.  
ACCESSION M29774 M29774.1 GI:178176  
VERSION 1  
KEYWORDS ADP-D-ribosyltransferase; NAD+ ADP-ribosyltransferase.  
SEGMENT 11 of 23  
SOURCE Human HeLa cell line, cDNA to mRNA, clones Hz[19,20,21,22,23,24].  
and liver DNA.  
ORGANISM Homo sapiens  
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;  
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.  
REFERENCE 1 (bases 1 to 20; 70 to 89)  
AUTHORS Auer,B., Nagl,U., Herzog,H., Schneider,R. and Schweiger,M.  
TITLE Human nuclear NAD+ ADP-ribosyltransferase (polymerizing):  
organization of the gene  
JOURNAL DNA 8 (8), 575-580 (1989)  
MEDLINE 90091744  
PUBMED 2513174  
REFERENCE 2 (bases 11 to 79)  
AUTHORS Herzog,H., Zabel,B.U., Schneider,R., Auer,B., Hirsch-Kauffmann,M.  
and Schweiger,M.  
TITLE Human nuclear NAD+ ADP-ribosyltransferase: localization of the gene  
on chromosome 1q41-q42 and expression of an active human enzyme in  
Escherichia coli  
JOURNAL Proc. Natl. Acad. Sci. U.S.A. 86 (10), 3514-3518 (1989)  
MEDLINE 89284454  
PUBMED 2498872  
COMMENT Draft entry and computer-readable sequence for [2] kindly submitted  
by H.Herzog, 07-MAR-1989.  
FEATURES  
Location/Qualifiers  
Source 1..89  
/organism="Homo sapiens"  
/db\_xref="taxon:9606"  
/map="1q41-q42"  
intron <1..10  
/gene="PPOL"  
exon /note="PPOL intron J"  
11..79  
/gene="PPOL"  
/note="NAD+ ADP-ribosyltransferase, (EC 2.4.2.30; 5' end  
put.); G00-119-508; putative"  
/number=11  
intron 80..>89  
/gene="PPOL"  
/note="PPOL intron K"  
BASE COUNT 33 a 17 c 19 g 20 t  
ORIGIN About 0.6 kb after segment 10; chromosome 1q41-q42.  
Query Match 68.0%; Score 13.6; DB 9; Length 89;  
Best Local Similarity 80.0%; Pred. No. 8.le+04;  
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;  
OY 1 AAGCGAAATGAATTGACT 20  
II II IIIIIIIII III  
Db 26 AAAAGAGATGAATTAAC 45  
Search completed: December 12, 2002, 02:56:19  
Job time : 323.116 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:04:50 ; Search time 92.087 Seconds  
(without alignments)  
440.192 Million cell updates/sec

Title: US-09-355-254F-9  
Perfect score: 18  
Sequence: 1 ggaatgacgttcctcgtg 18  
Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues  
Total number of hits satisfying chosen parameters: 2390332

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : N\_Geneseq\_101002:\*

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3:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1982.DAT:*
4:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1983.DAT:*
5:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1984.DAT:*
6:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1985.DAT:*
7:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1986.DAT:*
8:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1987.DAT:*
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19:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1998.DAT:*
20:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA1999.DAT:*
21:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2000.DAT:*
22:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2001A.DAT:*
23:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2001B.DAT:*
24:	/SIDS2/gcgdata/geneseq/geneseqn-emb1/NA2002.DAT:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	18	100.0	18	AAV46001	Immune adjuvant IL
2	18	100.0	18	AAV46001	Immune adjuvant IL
3	14.4	80.0	65	AAV46001	Murine Toll-like r
4	13.8	76.7	38	AAV46001	Rat spliced transc
5	13.8	76.7	89	AAV46001	Proteoglycan core
6	13.8	76.7	91	AAV46001	Bacillus clausii g
7	13.4	74.4	60	AAV46001	Human secreted pro
8	13.2	73.3	65	AAV46001	Human spliced tran
9	13.2	73.3	65	AAV46001	Rat spliced transc

10	13.2	73.3	65	24	ABN58058	Mouse spliced tra
11	13.2	73.3	90	22	AAK22893	Human brain expres
12	13.2	73.3	90	22	AAK49067	Human bone marrow
13	13.2	73.3	90	22	AAI54894	Probe #23580 used
14	13	72.2	28	19	AAV23036	HG3305S-28 primer
15	12.8	71.1	29	22	AAV74984	E. nidulans PPyrG
16	12.8	71.1	50	9	AAAB1197	Pertussis S2 subun
17	12.8	71.1	65	24	ABN29052	Rat spliced transc
18	12.8	71.1	96	16	AAI19415	Human gene signat
19	12.4	68.9	22	21	AAI287493	Human ADH7 gene ex
20	12.4	68.9	26	24	AAI32654	Human ACE-2 DNA fr
21	12.4	68.9	51	22	AAH79539	Human DNA containi
22	12.4	68.9	60	24	ABN49023	Human spliced tran
23	12.4	68.9	70	17	AAI79765	Nucleic acid inhib
24	12.4	68.9	70	21	AAI252674	High salt SELEX hu
25	12.4	68.9	72	16	AAI21588	Human gene signat
26	12.4	68.9	85	24	ABK15932	Human lung tumour
27	12.2	67.8	24	19	AAV09031	Human hhl sodium c
28	12.2	67.8	30	24	ABK11663	Rabbit calreticuli
29	12.2	67.8	33	15	AAO63125	Extension motif fo
30	12.2	67.8	41	24	ABL50365	Human ribosomal pr
31	12.2	67.8	41	24	ABL50366	Human ribosomal pr
32	12.2	67.8	50	23	ABL01010	Human SNP involvin
33	12.2	67.8	51	23	ABL00204	Human silent nonco
34	12.2	67.8	65	24	ABN28425	Rat spliced transc
35	12	66.7	33	19	AAV54224	Murine T-cell rece
36	12	66.7	33	19	AAV54228	Primer KC117 used
37	12	66.7	33	20	AAV55310	Soluble sc-TCR fus
38	12	66.7	33	20	AAV55306	Soluble sc-TCR fus
39	12	66.7	56	16	AAO86442	FIPV SM protein DN
40	12	66.7	56	22	AAH43094	PCR primer used to
41	12	66.7	77	22	ABA70510	Human foetal liver
42	12	66.7	77	22	ABA37136	Probe #15602 for g
43	12	66.7	77	22	AAK18755	Human brain expres
44	12	66.7	77	22	AAK44891	Human bone marrow
45	12	66.7	77	22	AAI24978	Probe #14911 for g

ALIGNMENTS

RESULT 1  
AAV46001  
ID AAV46001 standard; DNA; 18 BP.

AC AAV46001;

XX 16-OCT-1998 (first entry)

DT Immune adjuvant IL-13.

XX Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;

XX modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;

XX Ig class; autoimmune response; T-cell; B-cell; tumour; ss.

XX Class Bacteria.

XX EP855184-A1.

XX 29-JUL-1998.

XX 23-JAN-1997; 97EP-0101019.

XX 23-JAN-1997; 97EP-0101019.

XX (HEG/) HEG K.

XX (LIPF/) LIPFORD G B.

XX (WAGN/) WAGNER H.

XX Heeg K, Lipford GB, Wagner H;

XX WPI; 1998-389630/34.

XX

PT Antigenic composition comprises polynucleotide fragment and antigen  
PT - used as vaccine to treat or prevent e.g. cancer or pathogen  
PT infections and to modulate immune response e.g. tolerance break and  
PT regulation of TH1/TH2 cells  
PS Example 5; Page 8; 28pp; English.  
XX  
XX AAV45993-V46019 are fragments of bacterial polynucleotides which are  
XX used as immune adjuvants for inclusion into vaccines to treat cancer and  
XX for prophylaxis and/or treatment of conditions caused by pathogenic  
XX micro-organisms. The polynucleotide is used for modulation of an immune  
XX response and the modulation is selected from the group break of  
XX tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig  
XX classes, treatment of autoimmune responses and induction of tolerances.  
XX DNA oligomers are used to enhance the reactivity of immune cells to  
XX viral, bacterial and parasitic antigens, to break tolerance in anergic T  
XX and B cells e.g. against tumour antigens, as adjuvants in vaccination  
XX against tumour defined antigens and immunostimulatory substances in an  
XX immune response against tumours and to suppress immune reactions of the  
XX innate and acquired immune system. The composition is inexpensive and  
XX stable and does not cause lethal shock, which happens with prior art  
XX bacterial sequences.  
SQ Sequence 18 BP; 3 A; 4 C; 6 G; 5 T; 0 other;  
Query Match 100.0%; Score 18; DB 19; Length 18;  
Best Local Similarity 100.0%; Pred. No. 2.3;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGAATGACGTTCCCTGTG 18  
Db 1 GGAATGACGTTCCCTGTG 18  
RESULT 2  
AAL39185  
ID AAL39185 standard; DNA; 18 BP.  
XX  
XX AAL39185;  
XX  
XX  
XX 05-SEP-2002 (first entry)  
XX Murine Toll-like receptor related CpG DNA SEQ ID NO 60.  
XX Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.  
XX Unidentified.  
XX WO200222809-A2.  
XX  
XX 21-MAR-2002.  
XX  
XX 17-SEP-2001; 2001WO-US29229.  
XX  
XX 15-SEP-2000; 2000US-233035P.  
XX 23-JAN-2001; 2001US-263657P.  
XX 17-MAY-2001; 2001US-291726P.  
XX 22-JUN-2001; 2001US-300210P.  
XX  
XX (COLE-) COLEY PHARM GMBH.  
XX Bauer S., Lipford G., Wagner H;  
XX WPI; 2002-393964/42.  
XX  
XX New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,  
XX useful for identifying species specificity of immunostimulatory nucleic  
XX acid and identifying immunostimulatory nucleic acids  
XX  
XX Disclosure; Page 76; 195pp; English.  
XX  
XX The invention relates to isolated murine Toll-like receptors (TLR)9,  
XX TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined

CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or  
CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their  
CC fragments have an amino acid sequence which is identical to human TLR9,  
CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino  
CC acid of a murine TLR polypeptide. The isolated nucleic acids of the  
CC invention are useful for inhibiting TLR9 signalling activity in a cell.  
CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid  
CC molecules which interact with a TLR polypeptide or its fragment. The  
CC TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The  
CC TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9  
CC signalling activity of a test compound (that is not a nucleic acid, and  
CC is a polypeptide or a part of a combinatorial library of compounds) with  
CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for  
CC identifying species specificity of an ISNA. The isolated nucleic acids of  
CC the invention are useful as probes or primers. This polynucleotide  
CC sequence represents DNA relating to the isolated toll-like receptors of  
CC the invention.  
SQ Sequence 18 BP; 3 A; 4 C; 6 G; 5 T; 0 other;  
Query Match 100.0%; Score 18; DB 24; Length 18;  
Best Local Similarity 100.0%; Pred. No. 2.3;  
Matches 18; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GGAATGACGTTCCCTGTG 18  
Db 1 GGAATGACGTTCCCTGTG 18  
RESULT 3  
ABN30296  
ID ABN30296 standard; DNA; 65 BP.  
XX  
XX ABN30296;  
XX  
XX 15-JUL-2002 (first entry)  
XX Rat spliced transcript detection oligonucleotide SEQ ID NO:3044.  
XX Human; mouse; rat; splice transcript; detection; RNA transcript;  
XX splice variant; transcriptome; oligonucleotide library; ss.  
XX Rattus norvegicus.  
XX  
XX WO200210449-A2.  
XX  
XX 07-FEB-2002.  
XX  
XX 20-JUL-2001; 2001WO-IB01903.  
XX  
XX 28-JUL-2000; 2000US-221607P.  
XX 02-MAY-2001; 2001US-287724P.  
XX  
XX (COMP-) COMPUGEN INC.  
XX Shoshan A., Wasserman A., Mintz E., Mintz L., Faigler S;  
XX WPI; 2002-257383/30.  
XX  
XX New oligonucleotide libraries comprising oligonucleotides which  
XX selectively hybridize to mRNAs transcribed from a transcription unit of  
XX a genome, useful for detecting tissue-, pathology-, and  
XX developmental-specific genes  
XX  
XX Example 1; SEQ ID 3044; 47pp; English.  
XX The present invention describes oligonucleotide libraries for detecting  
XX messenger RNAs that populate a (sub-)transcriptome, where the  
XX (sub-)transcriptome comprises messenger RNAs transcribed from multiple  
XX transcription units that populate a genome. The library comprises  
XX several oligonucleotides, each capable of hybridising selectively to a  
XX set of messenger RNAs transcribed from a given transcription unit of  
XX the genome, which encodes one or more messenger RNA splice variants.

CC The oligonucleotide libraries are useful for detecting mRNAs from a  
 CC biological sample, in expression profiling studies, in qualitatively or  
 CC quantitatively characterising the corresponding transcriptome, and in  
 CC detecting RNA transcripts and splice variants of human or animal  
 CC transcripts. The libraries may also be used as specialised mini  
 CC libraries to detect transcripts of a sub-transcriptome under a  
 CC particular biological or pathological state, and so allowing the  
 CC detection of tissue- and pathology-specific genes such as those genes  
 CC only expressed in specific tissue under a specific pathological  
 CC condition; to detect developmental specific genes; and to detect RNA  
 CC transcripts and splice variants of a transcriptome of a patient suffering  
 CC from a particular disorder. ABN27253 to ABN5589 represent  
 CC oligonucleotide sequences from rats, humans and mice, which are used in  
 CC the exemplification of the present invention.  
 CC N.B. The sequence data for this patent did not form part of the printed  
 CC specification, but was obtained in electronic format directly from WIPO  
 CC at ftp.wipo.int/pub/published\_pct\_sequences.

SQ Sequence 65 BP; 18 A; 13 C; 16 G; 18 T; 0 other;

Query Match 80.0%; Score 14.4; DB 24; Length 65;

Best Local Similarity 93.8%; Pred. No. 2.5e+02;

Matches 15; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 3 AATGACGTTCCCTGTG 18

|||||

Db 9 AATGACGTTCCCTGCG 24

RESULT 4

AAQ57712/C

ID AAQ57712 standard; DNA; 38 BP.

XX AC

XX AAQ57712;

XX 11-AUG-1994 (first entry)

DE Proteoglycan core glycoprotein antisense primer.

XX XX

KW Neurocan; cell adhesion; leukocyte-endothelial cell recognition; lipid;  
 KW tissue-related inflammation allergy; cellular; humoral; carbohydrate;  
 KW hypersensitivity; trauma; neuronal development; cell transport; enzyme;  
 KW infection; diagnosis; lectin; versican; aggrecan; gelsolin; saccharide;  
 KW receptor; cell recognition; membrane cytoplasmic protein; nucleoside;  
 KW ion; ss.

XX Synthetic.

XX XX

PH Key Location/Qualifiers

FT modified\_base 28

FT /\*tag= a

FT /label= Inosine

XX XX

PN WO9403601-A.

XX XX

PD 17-FEB-1994.

XX XX

PF 03-AUG-1993; 93WO-US07306.

XX XX

PR 03-AUG-1992; 92US-0922911.

XX XX

PA (UYNV ) UNIV NEW YORK STATE.

XX XX

PI Margolis RK, Margolis RU, Rauch U;

XX XX

DR WPI; 1994-065690/08.

XX XX

PT Eukaryotic neurocan polypeptide(s) with epidermal growth factor,  
 PT lectin or complement binding activity - used in the diagnosis,  
 PT treatment or research of hypersensitivity and allergic diseases  
 XX XX  
 PS Example 1; Fig 1B; 105pp; English.

XX

CC The sequences given in AAQ57711-12 represent primers which were used  
 CC for the isolation of the neurocan cDNA. The neurocan protein  
 CC has several biological activities, including cell adhesion, leukocyte-  
 CC endothelial cell recognition, tissue-related inflammation allergies,  
 CC cellular and/or humoral hypersensitivity, trauma, neuronal  
 CC development, and cell transport and/or infection. Compositions  
 CC containing them can be used as modulators of these conditions, and  
 CC may be used as therapeutic, diagnostic, and/or research tools.  
 CC Neurocan peptides can be used to mimic proteins, such as lectins,  
 CC cell adhesion molecules, versicans, aggrecans or gelsolins, as  
 CC receptor or effector subtypes. The protein can be used to treat  
 CC diseases involving a qualitative or quantitative pathological  
 CC abnormality of cell adhesion or leukocyte-endothelial cell recognition,  
 CC or a functionally associated molecule such as a membrane cytoplasmic  
 CC protein, lipid, carbohydrate, saccharide, nucleoside, enzyme or ion.  
 XX XX

SQ Sequence 38 BP; 7 A; 9 C; 5 G; 10 T; 7 other;

Query Match 76.7%; Score 13.8; DB 15; Length 38;

Best Local Similarity 70.6%; Pred. No. 5.2e+02;

Matches 12; Conservative 3; Mismatches 2; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGT 17

|||||

Db 38 GGAATGATGTCCTGT 22

RESULT 5

ABK79561/C

ID ABK79561 standard; DNA; 89 BP.

XX AC

XX ABK79561;

XX 13-AUG-2002 (first entry)

XX XX

DE Bacillus clausii genomic sequence tag (GST) #2404.

KW Differential gene expression; genomic sequenced tag; GST;  
 KW altered culture condition; environmental stress;  
 KW physiological provocation; ds.

XX Bacillus clausii.

XX WO200229113-A2.

XX 11-APR-2002.

XX XX

PF 05-OCT-2001; 2001WO-US31437.

XX XX

PR 06-OCT-2000; 2000US-0680598.

XX XX

PR 27-MAR-2001; 2001US-279526P.

XX XX

PA (NOVO ) NOVOZYMES BIOTECH INC.

XX XX

PI (NOVO ) NOVOZYMES AS.

XX XX

PI Berka R, Clausen IG;

XX XX

DR WPI; 2002-416684/44.

XX XX

PT Monitoring differential expression of several genes in first Bacillus  
 PT cell relative to expression of same genes in one or more second  
 PT Bacillus cells, by using substrate containing Bacillus genomic  
 PT sequenced tag array

PS Claim 11; SEQ ID NO 6852; 200pp; English.

XX XX

CC The invention describes a method of monitoring differential expression of  
 CC genes in a first Bacillus cell relative to expression of the genes in  
 CC other Bacillus cells, comprising hybridising labelled nucleic acid probes  
 CC isolated from Bacillus cells to a substrate containing array of Bacillus  
 CC genomic sequenced tags (GST), examining the array, and determining  
 CC relative gene expression by an observed hybridisation reporter signal of  
 CC a spot in the array. The method is useful for measuring the expression of





CC the exemplification of the present invention.  
CC N.B. The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX Sequence 65 BP; 15 A; 20 C; 19 G; 11 T; 0 other;

Query Match 73.3%; Score 13.2; DB 24; Length 65;  
Best Local Similarity 83.3%; Pred. No. 1.2e+03;  
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18  
||||| |  
Db 43 GGAAGGACGACCCCTGTG 60

RESULT 10  
ABN58058  
ID ABN58058 standard; DNA; 65 BP.  
XX  
AC ABN58058;  
XX  
DT 15-JUL-2002 (first entry)  
XX  
DE Mouse spliced transcript detection oligonucleotide SEQ ID NO:30806.

XX Human; mouse; rat; splice transcript; detection; RNA transcript;  
KW splice variant; transcriptome; oligonucleotide library; ss...

XX Mus musculus.

XX WO200210449-A2.

XX 07-FEB-2002.

XX 20-JUL-2001; 2001WO-IB01903.

XX 28-JUL-2000; 2000US-221607P.

XX 02-MAY-2001; 2001US-287724P.

XX (COMP-) COMPUEN INC.

XX Shoshan A, Wasserman A, Mintz E, Mintz L, Faigler S;

XX WPI; 2002-257383/30.

XX New oligonucleotide libraries comprising oligonucleotides which  
PT selectively hybridize to mRNAs transcribed from a transcription unit of  
PT a genome, useful for detecting tissue-, pathology-, and  
PT developmental-specific genes

XX Example 1; SEQ ID 30806; 47pp; English.

XX The present invention describes oligonucleotide libraries for detecting  
CC messenger RNAs that populate a (sub-)transcriptome, where the  
CC (sub-)transcriptome comprises messenger RNAs transcribed from multiple  
CC transcription units that populate a genome. The library comprises  
CC several oligonucleotides, each capable of hybridising selectively to a  
CC set of messenger RNAs transcribed from a given transcription unit of  
CC the genome, which encodes one or more messenger RNA splice variants.  
CC The oligonucleotide libraries are useful for detecting mRNAs from a  
CC biological sample, in expression profiling studies, in qualitatively or  
CC quantitatively characterising the corresponding transcriptome, and in  
CC detecting RNA transcripts and splice variants of human or animal  
CC transcriptomes. The libraries may also be used as specialised mini  
CC libraries to detect transcripts of a sub-transcriptome under a  
CC particular biological or pathological state, and so allowing the  
CC detection of tissue- and pathology-specific genes such as those genes  
CC only expressed in specific tissue under a specific pathological  
CC condition; to detect developmental specific genes; and to detect RNA  
CC transcripts and splice variants of a transcriptome of a patient suffering  
CC from a particular disorder. ABN27253 to ABN59589 represent  
CC oligonucleotide sequences from rats, humans and mice, which are used in

CC the exemplification of the present invention.  
CC N.B. The sequence data for this patent did not form part of the printed  
CC specification, but was obtained in electronic format directly from WIPO  
CC at ftp.wipo.int/pub/published\_pct\_sequences.

XX Sequence 65 BP; 20 A; 17 C; 15 G; 13 T; 0 other;

Query Match 73.3%; Score 13.2; DB 24; Length 65;  
Best Local Similarity 83.3%; Pred. No. 1.2e+03;  
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18  
||||| |  
Db 15 GGAATGAAGTTCCTGTG 32

RESULT 11  
AAK22893/c  
ID AAK22893 standard; DNA; 90 BP.  
XX  
AC AAK22893;  
XX  
DT 05-NOV-2001 (first entry)  
XX  
DE Human brain expressed single exon probe SEQ ID NO: 22884.

XX Human; brain expressed exon; gene expression analysis; probe;  
KW microarray; Alzheimer's disease; multiple sclerosis; schizophrenia;  
KW epilepsy; cancer; ss.

XX Homo sapiens.

XX WO200157275-A2.

XX 09-AUG-2001.

XX 30-JAN-2001; 2001WO-US00667.

XX 04-FEB-2000; 2000US-0180312.

XX 26-MAY-2000; 2000US-0207456.

XX 30-JUN-2000; 2000US-0608408.

XX 03-AUG-2000; 2000US-0632366.

XX 21-SEP-2000; 2000US-0234687.

XX 27-SEP-2000; 2000US-0236359.

XX 04-OCT-2000; 2000GB-0024263.

XX (MOLE-) MOLECULAR DYNAMICS INC.

XX Penn SG, Hanzel DK, Chen W, Rank DR;

XX WPI; 2001-483446/52.

XX Single exon nucleic acid probes for analyzing gene expression in human  
PT brains -

XX Example 4; SEQ ID NO: 22884; 650pp + Sequence Listing; English.

XX The present invention provides a number of single exon nucleic acid  
CC probes which are derived from genomic sequences expressed in the human  
CC brain. They can be used to measure gene expression in brain cell samples,  
CC which may enable the diagnosis and improved treatment of nervous system  
CC diseases such as Alzheimer's disease, multiple sclerosis, schizophrenia,  
CC epilepsy and cancers. The present sequence is one of the probes of the  
CC invention.

XX Sequence 90 BP; 18 A; 33 C; 15 G; 24 T; 0 other;

Query Match 73.3%; Score 13.2; DB 22; Length 90;  
Best Local Similarity 83.3%; Pred. No. 1.2e+03;  
Matches 15; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 GGAATGACGTTCCCTGTG 18  
||||| |





PT New hepatitis G virus gene - useful for diagnosing and treating  
 PT diseases caused by virus  
 XX  
 XX Disclosure; Page 6; 128pp; Japanese.  
 XX  
 CC PCR primers AAV23018-74 were used to amplify and isolate new Hepatitis g  
 CC virus gene (see AAV23075-83 for gene fragments). RNA was synthesised  
 CC from the serum of nine patients judged positive for Hepatitis g virus  
 CC and cDNA synthesised from this RNA. The cDNA was used as a template in  
 CC several PCR reactions to isolate fragments of the new gene. The gene  
 CC may be useful for diagnosing and developing treatments for Hepatitis g  
 CC virus diseases.  
 XX  
 XX  
 SQ Sequence 28 BP; 8 A; 8 C; 5 G; 7 T; 0 other;  
 Query Match 72.2%; Score 13; DB 19; Length 28;  
 Best Local Similarity 100.0%; Pred. No. 1.4e+03; Length 28;  
 Matches 13; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 4 ATGACGTTCCCTG 16  
 |||||  
 Db 28 ATGACGTTCCCTG 16

RESULT 15  
 AAF74984  
 ID AAF74984 standard; DNA; 29 BP.  
 XX  
 AC AAF74984;  
 XX  
 DT 23-MAY-2001 (first entry)  
 XX  
 DE E. nidulans PPyrG sense primer PPyr3'.  
 XX  
 KW GlacA; promoter; PPyrG; PkIA; LacA; terminator; PCR primer; adapter;  
 KW protein expression; filamentous fungus; Escherichia coli;  
 KW Aspergillus niger; expression vector; ss.  
 XX  
 OS Emericella nidulans.  
 XX  
 PN WO200120007-A1.  
 XX  
 PD 22-MAR-2001.  
 XX  
 PF 13-SEP-2000; 2000WO-CA01084.  
 XX  
 PR 13-SEP-1999; 99US-0153228.  
 XX  
 PA (UYCO-) UNIV CONCORDIA.  
 XX  
 PI Sillaots S, Martinez-Perez A, Tsang A, Storms R;  
 XX  
 DR WPI; 2001-244813/25.  
 XX

PT Expression vector for isolating filamentous fungi that expresses a  
 PT protein of interest at high levels, has selectable marker for fungi and  
 PT promoter operably linked to nucleic acid sequence encoding the protein  
 PT

Example 3; Page 33; 54pp; English.

XX The present invention describes an expression vector (I) capable of  
 CC enabling a systematic analysis of gene expression and/or high-throughput  
 CC strain improvement screens in filamentous fungi, comprising a selectable  
 CC marker for the filamentous fungi and a promoter operably linked to a  
 CC sequence encoding a protein of interest or its part. Also described is a  
 CC filamentous fungi strain (II), enabling the expression of a protein of  
 CC interest to greater than 100, preferably greater than 1000 times  
 CC compared to the level in a parent strain. (I) is useful for isolating a  
 CC filamentous fungi, in particular Aspergillus niger that expresses a  
 CC protein of interest at high levels required for industrial-scale protein  
 CC production. (I) increases the expression of the protein of interest by  
 CC 100 times, preferably 1000 times, as compared to the level of expression

CC in a control filamentous fungi strain. (I) increases the efficiency and  
 CC reduces the time required for isolating strains that express proteins of  
 CC interest at high levels. AAF74980 to AAF74997 represent oligonucleotide  
 CC sequences used in an example from the present invention.

XX  
 SQ Sequence 29 BP; 8 A; 8 C; 5 G; 8 T; 0 other;  
 Query Match 71.1%; Score 12.8; DB 22; Length 29;  
 Best Local Similarity 87.5%; Pred. No. 1.8e+03;  
 Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 2 GAATGACGTTCCCTGT 17  
 |||||  
 Db 2 GAATGACGTTCCCTTT 17

Search completed: December 12, 2002, 01:36:19  
 Job time : 96.087 secs

GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:06:16 ; Search time 318.116 Seconds  
(without alignments)  
1829.698 Million cell updates/sec

Title: US-09-355-254F-23

Perfect score: 20  
Sequence: 1 caggcataacggttcogtag 20

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database :

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GenEmbl:
1: gb_da:*
2: gb_htg:*
3: gb_in:*
4: gb_om:*
5: gb_ov:*
6: gb_pat:*
7: gb_ph:*
8: gb_pl:*
9: gb_pr:*
10: gb_ro:*
11: gb_sts:*
12: gb_sy:*
13: gb_un:*
14: gb_vi:*
15: em_ba:*
16: em_fun:*
17: em_hum:*
18: em_in:*
19: em_mu:*
20: em_om:*
21: em_or:*
22: em_ov:*
23: em_pat:*
24: em_ph:*
25: em_pi:*
26: em_ro:*
27: em_sts:*
28: em_un:*
29: em_vi:*
30: em_htg_hum:*
31: em_htg_inv:*
32: em_htg_other:*
33: em_htg_mus:*
34: em_htg_pln:*
35: em_htg_rod:*
36: em_htg_mam:*
37: em_htg_vrt:*
38: em_sy:*
39: em_htgo_hum:*
40: em_htgo_mus:*
41: em_htgo_other:*

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score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	20	100.0	20	6	A9802	Sequence 24
2	20	100.0	20	6	A90889	Sequence 24
3	20	100.0	20	6	AX455564	Sequence
4	15.2	76.0	25	6	AX196810	Sequence
5	13.6	68.0	24	6	AX445605	Sequence
6	13.4	67.0	31	6	BD002472	Gene comp
7	13	65.0	20	6	E07207	PCR primer
8	12.8	64.0	33	6	AX262268	Sequence
9	12.8	64.0	35	6	AX262303	Sequence
10	12.8	64.0	39	6	AX262285	Sequence
11	12.8	64.0	65	6	AX486035	Sequence
12	12.8	64.0	97	6	I44826	Sequence 1
13	12.6	63.0	33	6	A32217	Synthetic m
14	12.6	63.0	50	6	AX103395	Sequence
15	12.2	61.0	30	6	AX374961	Sequence
16	12.2	61.0	31	6	AR106260	Sequence
17	12.2	61.0	31	6	E61282	Method for
18	12.2	61.0	37	6	AR145069	Sequence
19	12.2	61.0	37	6	AR168079	Sequence
20	12.2	61.0	37	6	AR169792	Sequence
21	12.2	61.0	37	6	AR204850	Sequence
22	12	60.0	33	6	A48801	Sequence 2
23	12	60.0	36	6	AX247485	Sequence
24	12	60.0	39	6	AX052711	Sequence
25	12	60.0	51	6	AX159959	Sequence
26	12	60.0	63	6	AX482092	Sequence
27	12	60.0	72	9	HSU91302	Sequence
28	12	60.0	90	9	HSC1CHX13	Homo sapien
29	11.8	59.0	24	6	AX445387	Sequence
30	11.8	59.0	26	6	AR125091	Sequence
31	11.8	59.0	31	6	BD002473	Gene comp
32	11.8	59.0	33	6	AR008899	Sequence
33	11.8	59.0	33	6	AR087594	Sequence
34	11.8	59.0	35	6	A84225	Sequence 4
35	11.8	59.0	38	6	AR038218	Sequence
36	11.8	59.0	39	6	AX080158	Sequence
37	11.8	59.0	61	6	AX080156	Sequence
38	11.8	59.0	65	6	AX484924	Sequence
39	11.8	59.0	100	14	S82444	{3' region,
40	11.8	59.0	100	14	S82445	{3' region,
41	11.8	59.0	100	14	S82446	{3' region,
42	11.6	58.0	24	6	AX443631	Sequence
43	11.6	58.0	25	6	AX447613	Sequence
44	11.6	58.0	28	6	AR160111	Sequence
45	11.6	58.0	35	6	AR084174	Sequence

# ALIGNMENTS

RESULT 1  
A9802  
LOCUS  
DEFINITION  
ACCESSION  
VERSION  
KEYWORDS  
SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL

A9802  
Sequence 24 from Patent WO9832462.  
A9802  
A9802.1 GI:6738316  
unidentified.  
unidentified.  
unclassified.  
1 (bases 1 to 20)  
Lipford,G.B. and Heeg,K.  
PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
Patent: WO 9832462-A 24 30-JUL-1998;

20 bp  
DNA  
linear  
PAT 22-JAN-2000

Pred. No. is the number of results predicted by chance to have a

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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
  Location/Qualifiers
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      /organism="unidentified"
      /db_xref="taxon:32644"
BASE COUNT      5 a      5 c      6 g      4 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 9.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTAG 20
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Db 1 CAGGCATACGGTTCGGTAG 20

RESULT 2
LOCUS      A90889
DEFINITION Sequence 24 from Patent EP0855184.
ACCESSION A90889
VERSION A90889.1 GI:6739343
KEYWORDS   .
SOURCE     .
ORGANISM   .
REFERENCE  1 (bases 1 to 20)
AUTHORS    Heeg,K.P. and Lipford,G.B.
TITLE      Pharmaceutical composition comprising a polynucleotide and an
            antigen especially for vaccination
JOURNAL    Patent: EP 0855184-A 24 29-JUL-1998;
            HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
FEATURES   .
SOURCE     1..20
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BASE COUNT      5 a      5 c      6 g      4 t
ORIGIN

Query Match      100.0%; Score 20; DB 6; Length 20;
Best Local Similarity 100.0%; Pred. No. 9.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTAG 20
    |||||
Db 1 CAGGCATACGGTTCGGTAG 20

RESULT 3
LOCUS      AX455564
DEFINITION Sequence 41 from Patent WO222809.
ACCESSION AX455564
VERSION AX455564.1 GI:21714632
KEYWORDS   .
SOURCE     .
ORGANISM   .
REFERENCE  1
AUTHORS    Bauer,S., Lipford,G. and Wagner,H.
TITLE      Process for high throughput screening of cpq-based
            immuno-agonist/antagonist
JOURNAL    Patent: WO 0222809-A 41 21-MAR-2002;
            Coley Pharmaceutical GmbH (DE)
FEATURES   .
SOURCE     1..20
            /organism="synthetic construct"
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            /note="Synthetic oligonucleotide"
BASE COUNT      5 a      5 c      6 g      4 t
ORIGIN

LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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Best Local Similarity 100.0%; Pred. No. 9.7;
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTAG 20
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Db 1 CAGGCATACGGTTCGGTAG 20

RESULT 4
LOCUS      AX196810
DEFINITION Sequence 517 from Patent WO0151627.
ACCESSION AX196810
VERSION AX196810.1 GI:15387016
KEYWORDS   .
SOURCE     .
ORGANISM   .
REFERENCE  1 (bases 1 to 25)
AUTHORS    Haug, B.M., Wang, M.L., Parsons, J.D. and Parnell, L.D.
TITLE      Nucleic acid molecules and other molecules associated with soybean
            cyst nematode resistance
JOURNAL    Patent: WO 0151627-A 517 19-JUL-2001;
            MONSANTO COMPANY (US)
FEATURES   .
SOURCE     1..25
            /organism="Glycine max"
            /db_xref="taxon:3847"
            /note="Seq ID: 240017_region_G3_6395_12_Forward_Primer"
BASE COUNT      6 a      6 c      6 g      7 t
ORIGIN

Query Match      76.0%; Score 15.2; DB 6; Length 25;
Best Local Similarity 85.0%; Pred. No. 3.6e+03;
Matches 17; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTAG 20
    |||||
Db 22 CAGGCACCGGTTTCAGTAG 3

RESULT 5
LOCUS      AX445605
DEFINITION Sequence 2060 from Patent WO0216649.
ACCESSION AX445605
VERSION AX445605.1 GI:21692886
KEYWORDS   .
SOURCE     .
ORGANISM   .
REFERENCE  1
AUTHORS    Gunderson, K.
TITLE      Probes and decoder oligonucleotides
JOURNAL    Patent: WO 0216649-A 2060 28-FEB-2002;
            Illumina, Inc. (US)
FEATURES   .
SOURCE     1..24
            /organism="synthetic construct"
            /db_xref="taxon:32630"
            /note="Computer Generated Probe Sequence."
BASE COUNT      6 a      5 c      9 g      4 t
ORIGIN

Query Match      68.0%; Score 13.6; DB 6; Length 24;
Best Local Similarity 80.0%; Pred. No. 2.6e+04;
Matches 16; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTAG 20
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```

Db      2  CAGGTCATGACAGTTCGGTAG 21
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RESULT 6
BD002472
LOCUS   31 bp      DNA      linear      PAT 31-JAN-2002
DEFINITION
Gene composition and method.
ACCESSION
BD002472
VERSION 1 GI:18630433
KEYWORDS
JP 2000245487-A/138.
SOURCE  unidentified.
ORGANISM
unclassified.
REFERENCE
1 (bases 1 to 31)
AUTHORS
Sha,N., Walinton,J. and Patel,N.
TITLE
Gene composition and method
JOURNAL
Patent: JP 2000245487-A 138 12-SEP-2000;
AFIMETRICS INC
COMMENT
OS Unknown
PN JP 2000245487-A/138
PD 12-SEP-2000
PF 27-JAN-2000 JP 2000019392
PR 27-JAN-1999 US 09/238.402
PI NIRA SHA,JANET WALINTON,NIRA PATEL
PC C12N15/09,C12Q1/68,C12N15/00
CC
FH Key
FT source
FT Location/Qualifiers
FT /Organism='Unknown'.
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Location/Qualifiers
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BASE COUNT 10 a 6 c 5 g 9 t 1 others
ORIGIN
Query Match 67.0%; Score 13.4; DB 6; Length 31;
Best Local Similarity 82.4%; Pred. NO. 3.3e+04;
Matches 14; Conservative 1; Mismatches 2; Indels 0; Gaps 0;
QY 1 CAGGCATACAGTTCGG 17
|||||  ||  |||
Db      15  CRGGCAAACTGTTCGG 31
|||||  ||  |||
RESULT 7
E07207/c
LOCUS   20 bp      DNA      linear      PAT 29-SEP-1997
DEFINITION
PCR primer to detect Acholeplasma sp.
ACCESSION
E07207
VERSION 1 GI:2175348
KEYWORDS
JP 1994098800-A/2.
SOURCE  unidentified.
ORGANISM
unclassified.
REFERENCE
1 (bases 1 to 20)
AUTHORS
Nakagawa,T., Demori,T., Asada,K., Katou,I. and Harasawa,A.
TITLE
METHOD FOR DETECTING ACHOLEPLASMA
JOURNAL
Patent: JP 1994098800-A 2 12-APR-1994;
TAKARA SHUZO CO LTD
COMMENT
OS None
OC Artificial sequences.
PN JP 1994098800-A/2
PD 12-APR-1994
PF 21-SEP-1992 JP 1992274830
PI NAKAGAWA TOMOKO, UEMORI TAKASHI, ASADA KIYOZOU, PI KATO
IKUNOSHIN,
PI HARASAWA AKIRA
PC C12Q1/68,C12N15/11,C12Q1/04,(C12Q1/04,C12R1:01); CC
strandedness: Single;
CC topology: Linear;
FH Key
FT source
FT Location/Qualifiers

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FH      source
FT      /Organism='Artificial sequences'.
FEATURES
source
Location/Qualifiers
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BASE COUNT 4 a 7 c 3 g 4 t 2 others
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Best Local Similarity 76.5%; Pred. NO. 5.4e+04;
Matches 13; Conservative 2; Mismatches 2; Indels 0; Gaps 0;
QY 4 GCATAACGGTTCGGTAG 20
|||||  ||  |||
Db      18  GGATCAGGTTCSGTAR 2
|||||  ||  |||
RESULT 8
AX262268
LOCUS   33 bp      DNA      linear      PAT 26-OCT-2001
DEFINITION
Sequence 24 from Patent WO0173052.
ACCESSION
AX262268
VERSION 1 GI:16511217
KEYWORDS
synthetic construct.
SOURCE  synthetic construct.
ORGANISM
artificial sequences.
REFERENCE
1
AUTHORS
Mchenry,C.S.
TITLE
Thermophilic polymerase III holoenzyme
JOURNAL
Patent: WO 0173052-A 24 04-OCT-2001;
Mchenry, Charles S. (US)
FEATURES
source
Location/Qualifiers
1..33
/organism="synthetic construct"
/db_xref="taxon:32630"
/Note="ATG forward primer P118-S85"
BASE COUNT 13 a 10 c 5 g 5 t
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Query Match 64.0%; Score 12.8; DB 6; Length 33;
Best Local Similarity 87.5%; Pred. NO. 6.8e+04;
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 5 CATAACGGTTCGGTAG 20
|||||  ||  |||
Db      11  CATAACGGTTCCTCAAG 26
|||||  ||  |||
RESULT 9
AX262303
LOCUS   35 bp      DNA      linear      PAT 26-OCT-2001
DEFINITION
Sequence 59 from Patent WO0173052.
ACCESSION
AX262303
VERSION 1 GI:16511248
KEYWORDS
synthetic construct.
SOURCE  synthetic construct.
ORGANISM
artificial sequences.
REFERENCE
1
AUTHORS
Mchenry,C.S.
TITLE
Thermophilic polymerase III holoenzyme
JOURNAL
Patent: WO 0173052-A 59 04-OCT-2001;
Mchenry, Charles S. (US)
FEATURES
source
Location/Qualifiers
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/Note="forward/sense ATG primer P118-S78cia2"
BASE COUNT 15 a 7 c 6 g 7 t
ORIGIN

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Query Match 64.0%; Score 12.8; DB 6; Length 35;  
Best Local Similarity 87.5%; Pred. No. 6.8e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 CATAACGGTTCGGTAG 20  
IIIIIIIIIIIIIIII  
DB 17 CATAACGGTTCACAG 32

RESULT 10  
AX262285  
LOCUS AX262285 39 bp DNA linear PAT 26-OCT-2001  
DEFINITION Sequence 41 from Patent WO0173052.  
ACCESSION AX262285  
VERSION AX262285.1 GI:16511230  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.

REFERENCE 1  
AUTHORS McHenry, C.S.  
TITLE Thermophilic polymerase III holoenzyme  
JOURNAL Patent: WO 01/73052-A 41 04-OCT-2001;  
McHenry, Charles S. (US)  
FEATURES  
Location/Qualifiers  
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/db\_xref="taxon:32630"  
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15 a 9 c 7 g 8 t

BASE COUNT 15 a 9 c 7 g 8 t  
ORIGIN

Query Match 64.0%; Score 12.8; DB 6; Length 39;  
Best Local Similarity 87.5%; Pred. No. 6.8e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 5 CATAACGGTTCGGTAG 20  
IIIIIIIIIIIIIIII  
DB 21 CATAACGGTTCACAG 36

RESULT 11  
AX486035/c  
LOCUS AX486035 65 bp DNA linear PAT 16-AUG-2002  
DEFINITION Sequence 3335 from Patent WO02053728.  
ACCESSION AX486035  
VERSION AX486035.1 GI:22320251  
KEYWORDS  
SOURCE Candida albicans.  
ORGANISM Candida albicans

REFERENCE 1  
AUTHORS Roemer, T., Jiang, B., Boone, C., Bussey, H. and Ohlson, K.L.  
TITLE Gene disruption methodologies for drug target discovery  
JOURNAL Patent: WO 02053728-A 3335 11-JUL-2002;  
Eli Lilly Pharmaceuticals, Inc. (US)  
FEATURES  
Location/Qualifiers  
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/db\_xref="taxon:5476"  
18 a 10 c 11 g 26 t

BASE COUNT 18 a 10 c 11 g 26 t  
ORIGIN

Query Match 64.0%; Score 12.8; DB 6; Length 65;  
Best Local Similarity 87.5%; Pred. No. 6.7e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTC 16  
IIIIIIIIIIIIIIII  
DB 61 CAGTCATACGGATCC 46

RESULT 12  
I44826/c  
LOCUS I44826 97 bp DNA linear PAT 07-OCT-1997  
DEFINITION Sequence 1 from patent US 5635601.  
ACCESSION I44826  
VERSION I44826.1 GI:2469539  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
Unclassified.

REFERENCE 1 (bases 1 to 97)  
AUTHORS Moyle, M. and McLean, J.W.  
TITLE Beta-8 integrin subunit antibodies  
JOURNAL Patent: US 5635601-A 1 03-JUN-1997;  
FEATURES  
Location/Qualifiers  
1..97  
/organism="unknown"  
26 a 18 c 33 g 20 t

Query Match 64.0%; Score 12.8; DB 6; Length 97;  
Best Local Similarity 87.5%; Pred. No. 6.7e+04;  
Matches 14; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 3 GGCATAACGGTTCGGT 18  
IIIIIIIIIIIIIIII  
DB 30 GGCATCTCGGTTCCGT 15

RESULT 13  
A32217/c  
LOCUS A32217 33 bp DNA linear PAT 03-JUL-1996  
DEFINITION Synthetic mutagenic primer.  
ACCESSION A32217  
VERSION A32217.1 GI:1567327  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct  
artificial sequences.

FEATURES  
Location/Qualifiers  
1..33  
/organism="synthetic construct"  
/db\_xref="taxon:32630"  
5 a 10 c 7 g 11 t

BASE COUNT 5 a 10 c 7 g 11 t  
ORIGIN

Query Match 63.0%; Score 12.6; DB 6; Length 33;  
Best Local Similarity 78.9%; Pred. No. 8.8e+04;  
Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

QY 1 CAGGCATACGGTTCGGTA 19  
IIIIIIIIIIIIIIII  
DB 28 CAGAAGTAACGGATCCGTA 10

RESULT 14  
AX103395/c  
LOCUS AX103395 50 bp DNA linear PAT 30-MAY-2001  
DEFINITION Sequence 17 from Patent EP1104813.  
ACCESSION AX103395  
VERSION AX103395.1 GI:13919680  
KEYWORDS  
SOURCE unidentified adenovirus.  
ORGANISM unidentified adenovirus

REFERENCE 1 (bases 1 to 50)  
AUTHORS Rademaker, H.J., Fallaux, F.J., de Jong, R.N., van der Vliet, P.C. and  
Hoeben, R.C.  
TITLE Conditional replication of recombinant human adenovirus dna  
JOURNAL carrying modified inverted terminal repeat sequences  
Patent: EP 1104813-A 17 06-JUN-2001;  
Leids Universitair Medisch Centrum (NL)

```

FEATURES
  source      Location/Qualifiers
  1..50
    /organism="unidentified adenovirus"
    /db_xref="taxon:10535"
  misc_feature 1..50
    /note="BAD"
  BASE COUNT  15 a 10 c 10 g 15 t
  ORIGIN
    Query Match      63.0%; Score 12.6; DB 6; Length 50;
    Best Local Similarity 78.9%; Pred. No. 8.7e+04;
    Matches 15; Conservative 0; Mismatches 4; Indels 0; Gaps 0;

  QY 1 CAGGCATAACGGTTCCGTA 19
      || ||| || ||||| |||
  Db 33 CAGCCAAAGTGTTCGTA 15

  RESULT 15
  AX374961
  LOCUS      AX374961      30 bp      DNA      linear      PAT 01-MAR-2002
  DEFINITION Sequence 14 from Patent WO0210425.
  ACCESSION  AX374961
  VERSION    AX374961.1 GI:19169806
  KEYWORDS   .
  SOURCE     synthetic construct.
            ORIGIN      synthetic construct
            ORGANISM     artificial sequences.
  REFERENCE  1
  AUTHORS   Porto,D. and Sauer,M.
  TITLE     Ascorbic acid production from yeasts
  JOURNAL   Patent: WO 0210425-A 14 07-FEB-2002;
            Biopolo S.C.A.R.L. (IT)

  FEATURES
    source      Location/Qualifiers
    1..30
      /organism="synthetic construct"
      /db_xref="taxon:32630"
      /note="Forward PCR Primer for L-galactono-1,4-lactone
            dehydrogenase from A. thaliana"
    BASE COUNT      8 a 9 c 6 g 7 t
    ORIGIN

    Query Match      61.0%; Score 12.2; DB 6; Length 30;
    Best Local Similarity 82.4%; Pred. No. 1.4e+05;
    Matches 14; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

  QY 2 AGGCATAACGGTTCCGT 18
      ||||| ||| ||||| |||
  Db 6 AGCCATAAATGTTCGT 22

  Search completed: December 12, 2002, 02:56:25
  Job time : 324.116 secs

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GenCore version 5.1.3  
Copyright (c) 1993 - 2002 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:04:50 ; Search time 102.319 Seconds  
(without alignments)  
440.192 Million cell updates/sec

Title: US-09-355-254F-8

Perfect score: 20  
Sequence: 1 gattgcctgacgtcagagag 20

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2185239 seqs, 1125999159 residues

Total number of hits satisfying chosen parameters: 2390332

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

Database : N\_Geneseq\_101002.\*

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21: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2000.DAT.*
22: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001A.DAT.*
23: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2001B.DAT.*
24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	20	100.0	20	19	AAV46000
2	20	100.0	20	20	AAZ28189
3	20	100.0	20	21	AAZ29172
4	20	100.0	20	24	AAL39177
5	20	100.0	26	20	AAZ76047
6	20	100.0	27	18	AAT85832
7	20	100.0	27	20	AAV82454
8	20	100.0	27	20	AAV08336
9	20	100.0	27	22	AAI70581

10	20	100.0	27	22	AAH77396	Cyclic AMP respons
11	20	100.0	27	22	AAH77397	Cyclic AMP respons
12	20	100.0	27	22	AAF87956	Cyclic AMP respons
13	20	100.0	27	22	AAF87957	Cyclic AMP respons
14	20	100.0	27	22	AAF76267	CAMP response elem
15	20	100.0	27	24	ABA92274	CRE binding site o
16	20	100.0	27	24	ABA05538	Cyclic-AMP respons
17	19	95.0	20	19	AAV45997	Immune adjuvant CR
18	16.4	82.0	28	24	ABK14052	Cyclic AMP respons
19	16.4	82.0	32	22	AAF77813	CREB probe derived
20	16.4	82.0	37	19	AAV04084	Somatostatin gene
21	16.4	82.0	50	15	AAQ69701	Human somatostatin
22	16.4	82.0	50	18	AAI64163	Human somatostatin
23	16.4	82.0	50	20	AAI17451	Test sequence from
24	16.4	82.0	50	24	ABK82942	DNA binding molecu
25	15.8	79.0	20	19	AAV45999	Immune adjuvant CR
26	14.8	74.0	95	21	AAC30099	Human secreted pro
27	14.4	72.0	33	24	ABK87820	Somatostatin promo
28	14.4	72.0	33	24	ABK87821	Somatostatin promo
29	14.4	72.0	83	24	AA598555	Human Pleckstrin h
30	14.2	71.0	41	19	AAV50622	Brassica sp. polym
31	14.2	71.0	41	24	ABL96051	Brassica polymorph
32	14.2	71.0	60	24	ABN49177	Human spliced tran
33	13.8	69.0	65	24	ABN57982	Mouse spliced tran
34	13.6	68.0	51	22	AAI78809	Human silent SNP c
35	13.4	67.0	96	21	AAC12454	Human secreted pro
36	13.2	66.0	22	19	AAV04086	PEPCK gene CAMP re
37	13.2	66.0	25	22	AAH39807	SNP specific SNPE
38	13.2	66.0	47	21	AAZ68295	Human map-related
39	13.2	66.0	51	22	AAH39808	Human SNP flanking
40	13.2	66.0	60	24	ABN45698	Human spliced tran
41	13.2	66.0	62	22	AAI66152	Hepatitis E virus
42	13.2	66.0	65	24	ABN56672	Mouse spliced tran
43	13.2	66.0	90	24	ABK36416	HIV DNA encoding N
44	13	65.0	26	19	AAV46019	Immune adjuvant CR
45	12.8	64.0	21	19	AAV41019	Primer PL2FRARA:12

#### ALIGNMENTS

RESULT 1  
AAV46000  
ID AAV46000 standard; DNA; 20 BP.

XX  
AC AAV46000;

XX  
DT 16-OCT-1998 (first entry)

XX  
DE Immune adjuvant AP-1 #1.

XX  
KW Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;  
modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;  
KW Ig class; autoimmune response; T-cell; B-cell; tumour; ss.

XX  
OS Class Bacteria..

XX  
PN EP855184-A1.

XX  
PD 29-JUL-1998.

XX  
PF 23-JAN-1997; 97EP-0101019.

XX  
PR 23-JAN-1997; 97EP-0101019.

XX  
PA (HEG/) HEG K.

XX  
PA (LIPE/) LIPFORD G B.

XX  
PA (WAGN/) WAGNER H.

XX  
PI Heeg K, Lipford GB, Wagner H;

XX  
DR WPI; 1998-389630/34.

XX



CC when injected simultaneously. The peptides may also be used for  
 CC increasing inflammatory myocarditis in a mammal. Antibodies against the  
 CC peptides and the peptides themselves are used for measuring the risk of  
 CC inflammatory cardiomyopathy in a mammal. The peptides may also be used  
 CC in vaccines. Nucleic acids encoding the peptides may be used as  
 CC hybridization probes, e.g. in diagnostic assays to test for the  
 CC presence of Chlamydia DNA.

XX Sequence 20 BP; 5 A; 4 C; 7 G; 4 T; 0 other;  
 SQ Query Match 100.0%; Score 20; DB 21; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 0.48; 0; Indels 0; Gaps 0;  
 Matches 20; Conservative 0; Mismatches 0;

QY 1 GATTGCTGACGTCAGAGAG 20  
 Db 1 GATTGCTGACGTCAGAGAG 20  
 |||||

RESULT 4  
 AAL39177  
 ID AAL39177 standard; DNA; 20 BP.  
 XX  
 AC AAL39177;  
 XX  
 DT 05-SEP-2002 (first entry)  
 DE Murine Toll-like receptor related CpG DNA SEQ ID NO 52.  
 DE Murine Toll-like receptor; TLR8; TLR7; TLR8; ISNA; ds.  
 KW Unidentified.  
 XX  
 OS WO200222809-A2.  
 PN 21-MAR-2002.  
 XX  
 PD 17-SEP-2001; 2001WO-US29229.  
 XX  
 PF 15-SEP-2000; 2000US-233035P.  
 PR 23-JAN-2001; 2001US-263657P.  
 PR 17-MAY-2001; 2001US-291726P.  
 PR 22-JUN-2001; 2001US-300210P.  
 XX

PA (COLE-) COLEY PHARM GMBH.  
 XX  
 XX Bauer S, Lipford G, Wagner H;  
 XX WPI; 2002-393964/42.  
 DR  
 XX  
 XX New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,  
 PT useful for identifying species specificity of immunostimulatory nucleic  
 PT acid and identifying immunostimulatory nucleic acids  
 XX  
 PS Disclosure; Page 76; 195pp; English.  
 XX

CC The invention relates to isolated murine Toll-like receptors (TLR)9,  
 CC TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined  
 CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or  
 CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their  
 CC fragments have an amino acid sequence which is identical to human TLR9,  
 CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino  
 CC acid of a murine TLR polypeptide. The isolated nucleic acids of the  
 CC invention are useful for inhibiting TLR9 signalling activity in a cell.  
 CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid  
 CC molecules which interact with a TLR polypeptide or its fragment. The  
 CC TLR7, TLR8 or TLR9 polypeptides are also useful for identifying ISNA. The  
 CC TLR7, TLR8 and TLR9 polypeptides are also useful for comparing TLR9  
 CC signalling activity of a test compound (that is not a nucleic acid, and  
 CC is a polypeptide or a part of a combinatorial library of compounds) with  
 CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for  
 CC identifying species specificity of an ISNA. The isolated nucleic acids of  
 CC the invention are useful as probes or primers. This polynucleotide

CC sequence represents DNA relating to the isolated Toll-like receptors of  
 CC the invention.

SQ Sequence 20 BP; 5 A; 4 C; 7 G; 4 T; 0 other;

Query Match 100.0%; Score 20; DB 24; Length 20;  
 Best Local Similarity 100.0%; Pred. No. 0.48;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACGTCAGAGAG 20  
 Db 1 GATTGCTGACGTCAGAGAG 20  
 |||||

RESULT 5  
 AAX76047  
 ID AAX76047 standard; DNA; 26 BP.  
 XX  
 AC AAX76047;  
 XX  
 DT 30-JUL-1999 (first entry)  
 DE CAMP response element oligonucleotide SEQ ID NO:15.  
 DE  
 XX  
 KW CRE; cAMP response element; transcription factor decoy; cis-element;  
 KW tumour growth inhibitor; palindrome; hairpin; cancer; metabolism;  
 KW gene transcription regulation; inhibiting proliferation; ds.  
 XX  
 OS Synthetic.  
 XX  
 PN WO9926634-A1.  
 XX  
 PD 03-JUN-1999.  
 XX  
 PF 23-NOV-1998; 98WO-US25307.  
 XX  
 PR 24-NOV-1997; 97US-0977643.  
 XX  
 PA (USSH ) US DEPT HEALTH & HUMAN SERVICES.  
 XX  
 PI Cho-Chung YS;  
 XX  
 DR WPI; 1999-347612/29.  
 XX

PT Nucleic acids that compete with response elements for transcription  
 XX factors

PS Example 10; Page 54; 83pp; English.

CC The present invention describes a composition (A) comprising one or more  
 CC nucleic acids (I) that compete with cAMP (cyclic adenosine monophosphate)  
 CC response element (CRE) enhancer DNA for binding to transcription factors  
 CC (TF). (I) are used to regulate gene transcription in cells, in vitro or  
 CC in vivo, specifically for inhibiting proliferation of cancer cells, but  
 CC possibly also for regulation of metabolism in hepatitis B and other  
 CC viruses. HCT-15 human multidrug resistant colon carcinoma cells (2  
 CC million) were inoculated subcutaneously into the flank of nude mice,  
 CC then the CRE oligonucleotide 5'-TCAGCTTCATGCGTTCATGAGGTCA-3' injected  
 CC intraperitoneally at doses of 0.1 mg, 5 times per week, once the tumour  
 CC had reached 30-50 mg. This treatment resulted in over 85% reduction in  
 CC tumour growth, relative to an untreated control. (I) have high affinity  
 CC for TF and can inhibit growth of cancer cells without adverse effects on  
 CC normal cells (contrast use of antisense RNA). The method does not  
 CC require knowledge of the target gene sequence, only of the response  
 CC element sequence. The present sequence is used in the exemplification  
 CC of the present invention.

SQ Sequence 26 BP; 8 A; 4 C; 9 G; 5 T; 0 other;

Query Match 100.0%; Score 20; DB 20; Length 26;  
 Best Local Similarity 100.0%; Pred. No. 0.49;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACCTCAGAG 20  
 Db 4 GATTGCTGACCTCAGAG 23

## RESULT 6

AAAT85832  
 ID AAT85832 standard; DNA; 27 BP.

XX  
 AC AAT85832;

XX DT 21-NOV-1997 (first entry)

XX DE CRE oligonucleotide used in gel shift assay.

XX KW Activating transcription factor 1; ATF1; CREB; recognition sequence;  
 KW cyclic AMP responsive element binding protein; inhibition; binding;  
 KW proliferation; virus; cancer; HTLV1; leukaemia; antibody; ss.

XX OS Synthetic.

XX PN US5641486-A.

XX PD 24-JUN-1997.

XX PF 18-MAR-1994; 94US-0210880.

XX PR 18-MAR-1994; 94US-0210880.

XX PA (UYNE-) UNIV NEBRASKA.

XX PI Hinrichs SH, Orten DJ;

XX WPI; 1997-340900/31.

XX PT Inhibiting replication of cancer cells or viruses - with inhibitor  
 PT that binds to peptide sequence of activating transcription factor 1

XX PS Example 2; Column 6; 17pp; English.

XX This oligonucleotide sequence corresponds to the cyclic AMP binding  
 CC element (CRE) to which members of the activating transcription factor 1  
 CC (ATF1)-cyclic AMP responsive element binding protein (CREB) family  
 CC of protein bind. The sequence was used in a gel shift mobility assay to  
 CC identify agents which inhibit the binding of ATF1 to its recognition  
 CC sequence. The agents are preferably antibodies, small molecules or  
 CC polypeptides, especially the complementarity determining region of  
 CC monoclonal antibody Mab4. The agents cause inhibition of transcription  
 CC by dissociating ATF1 from its target gene and thus will prevent  
 CC proliferation of e.g. a virus or cancer cell, such as HTLV1-mediated  
 CC leukaemic cell proliferation.

XX SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 Other;

Query Match 100.0%; Score 20; DB 18; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.5;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACCTCAGAG 20  
 Db 4 GATTGCTGACCTCAGAG 23

## RESULT 7

AAV82454  
 ID AAV82454 standard; DNA; 27 BP.

XX AC AAV82454;

XX DT 12-APR-1999 (first entry)

XX DE ATF comp oligonucleotide used in competition analysis.

KW Vascular endothelial growth factor; VEGF; human; hypoxia;  
 KW vascular disease; tumour; cancer; angiogenesis; wound healing;  
 KW therapy; diagnosis; ds.

XX OS Synthetic.

XX OS Homo sapiens.

XX PN WO9856936-A1.

XX PD 17-DEC-1998.

XX PF 10-JUN-1998; 98WO-EP03517.

XX PR 10-JUN-1997; 97EP-0109418.

XX PA (PLAC ) MAX PLANCK GES FOERDERUNG WISSENSCHAFTEN.

XX PI Danert A, Plate K, Risau W;

XX WPI; 1999-080911/07.

XX PT New recombinant DNA - contains sequence that regulates

XX PT hypoxia-induced expression, used for, e.g. treatment and diagnosis  
 of vascular disease

XX PS Example 6; Page 41; 80pp; English.

XX CC Oligonucleotides hVEGF, hVEGF 5' DEL, AP1M1 and AP1M2 (see  
 CC AAV82449-52), and competitor oligonucleotides API comp, ATF comp  
 CC and VL30 (see AAV82453-55) were used in electrophoretic mobility  
 CC shift assays to determine which transcription factor(s) bind to  
 CC the cis-acting element that is involved in the potentiation of  
 CC hypoxia-inducible factor 1 (HIF-1) mediated hypoxic induction  
 CC of vascular endothelial growth factor (VEGF) gene regulatory  
 CC sequences. Experiments were performed using normoxic or hypoxic  
 CC C6 cell nuclear extracts. An API consensus binding site was shown  
 CC to compete for DNA-protein complex formation at potentiating  
 CC sequences. The invention relates to recombinant DNA molecules  
 CC comprising regulatory sequences of the VEGF gene, especially the  
 CC 3' untranslated region (see AAV82439) and promoter (see AAV82440),  
 CC being capable of modulating hypoxia inducible expression of a  
 CC heterologous DNA in vivo. Such recombinant DNA molecules, vectors,  
 CC host cells and transgenic animals can be used to identify and  
 CC develop compounds and methods for diagnosing, treating, preventing  
 CC and/or delaying a vascular or tumour disease.

XX SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 Other;

Query Match 100.0%; Score 20; DB 20; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.5;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCTGACCTCAGAG 20  
 Db 4 GATTGCTGACCTCAGAG 23

## RESULT 8

AAV08336

ID AAV08336 standard; DNA; 27 BP.

XX AC AAV08336;

XX DT 04-FEB-1999 (first entry)

XX DE CRE element coding sequence.

XX KW ATF1; activating transcription factor 1; inhibitor; gene transcription;  
 KW cell proliferation; cancer cell; human; ds.

XX OS Synthetic.

XX PN US5844096-A.

XX 01-DEC-1998.  
PD  
XX  
PF 20-DEC-1996; 96US-0771411.  
XX  
PR 18-MAR-1994; 94US-0210880.  
PR 20-DEC-1996; 96US-0771411.  
XX  
XX (UYNE-) UNIV NEBRASKA.  
PA  
XX Hinrichs SH, Orten DJ;  
PI  
XX  
XX  
DR WPI; 1999-044667/04.  
XX  
XX  
PT Inhibitor of activating transcription factor 1 mediated gene  
PT transcription - useful as anticancer or antiviral agent  
XX  
XX Example 2; Column 6; 17pp; English.  
XX  
XX This sequence represents a CRE element coding sequence. This sequence  
CC was used to test the effect of the inhibitory compound of the  
CC invention. The inhibitory compound binds to ATF1 residues 167-181 with  
CC sufficient affinity to dissociate ATF1 from a gene to which it is bound  
CC and thereby prevent transcription of the gene. The inhibitory compound  
CC and its derivatives are useful for inhibiting the ATF1-mediated  
CC proliferation of cancer cells and viruses, e.g. HTLV I.  
XX  
XX  
SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;  
Query Match 100.0%; Score 20; DB 20; Length 27;  
Best Local Similarity 100.0%; Pred. No. 0.5;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GATTGCTGACGTCAGAG 20  
DB 4 GATTGCTGACGTCAGAG 23  
|||||  
RESULT 9  
AAI70581  
ID AAI70581 standard; DNA; 27 BP.  
XX  
XX AAI70581;  
XX  
XX 21-JAN-2002 (first entry)  
XX  
XX Transcription factor CREB consensus oligonucleotide.  
DE  
XX  
XX Transcription factor; CREB; screening; detection; quantification;  
KW probe; ds.  
XX  
XX Synthetic.  
XX  
XX EP1136567-A1.  
PN  
XX  
XX 26-SEP-2001.  
PD  
XX  
XX 24-MAR-2000; 2000EP-0870057.  
PF  
XX  
XX 24-MAR-2000; 2000EP-0870057.  
PR  
XX  
XX (ADAR-) ADVANCED ARRAY TECHNOLOGIES SA.  
PA  
XX  
XX Remacle J, Renard P, Art M;  
PI  
XX  
XX WPI; 2001-640391/74.  
DR  
XX  
XX Screening, detecting or quantifying transcriptional factors in a  
PT biological sample comprises contacting the transcriptional factor with  
PT a double-stranded DNA sequence bound to an insoluble solid support -  
XX  
XX Example 4; Page 8; 20pp; English.  
PS  
XX

CC The present sequence is that of a CREB transcription factor  
CC consensus oligonucleotide. Double-stranded probe nucleotide  
CC sequences were constructed from 100 bp of a CMV 5' sequence (see  
CC AAI70578) linked to this oligonucleotide and used in microwell  
CC colorimetric CREB and phospho-CREB assays. The double-stranded  
CC probe was biotinylated at its CMV 5' extremity and linked to  
CC streptavidin-coated 96-wells plates. The plates were contacted  
CC with a nuclear extract of L929 murine fibrosarcoma cells,  
CC incubated with anti-CREB or anti-phospho-CREB antibody and then  
CC with peroxidase-conjugated antibody. The presence of CREB or  
CC phospho-CREB was detected through the action of peroxidase on  
CC tetramethylbenzidine. This is an example of the method of the  
CC invention, involving the detection of a transcription factor using  
CC a double-stranded DNA probe bound to an insoluble solid support at  
CC a concentration of at least 0.01 pmole/sq cm of support surface and  
CC at a distance of at least 6.8 nm from the surface of the support.  
CC The method allows the screening, detection and/or quantification of  
CC one or more transcriptional factors, of molecules binding such  
CC factors, and of molecules that inhibit such binding, using  
CC non-radioactive detection methods.  
XX  
SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;  
Query Match 100.0%; Score 20; DB 22; Length 27;  
Best Local Similarity 100.0%; Pred. No. 0.5;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 GATTGCTGACGTCAGAG 20  
DB 4 GATTGCTGACGTCAGAG 23  
|||||  
RESULT 10  
AAH77396  
ID AAH77396 standard; DNA; 27 BP.  
XX  
XX AAH77396;  
XX  
XX 05-NOV-2001 (first entry)  
XX  
XX Cyclic AMP response element CRE consensus oligonucleotide probe #1.  
DE  
XX  
XX Cancer; human; dithiocarbamate derivative; anticancer agent; probe; ss.  
KW  
XX  
XX Unidentified.  
XX  
XX US2001016600-A1.  
PN  
XX  
XX 23-AUG-2001.  
PD  
XX  
XX 12-DEC-2000; 2000US-0735205.  
PF  
XX  
XX 08-SEP-1998; 98US-0099390.  
PR  
XX 08-SEP-1999; 99US-0392122.  
PR  
XX 05-OCT-2000; 2000US-0679932.  
XX  
XX (KENN/) KENNEDY T P.  
PA  
XX  
XX Kennedy TP;  
PI  
XX  
XX WPI; 2001-557127/62.  
DR  
XX  
XX Treating cancer, asthma and cancer and reducing hypoxic or ischemic  
PT damage comprises administering dithiocarbamate thiolate anion or  
PT dithiocarbamate thiolate metal complex -  
XX  
XX Disclosure; Page 10; 36pp; English.  
PS  
XX  
XX The present invention describes a method of treating cancer, asthma and  
CC arthritis and reducing hypoxic or ischaemic damage, involving  
CC administering a dithiocarbamate thiolate anion or metal ion complex to  
CC the patient. The present sequence is a probe for the cyclic AMP response  
CC element CRE, which was described in the exemplification of the invention.  
CC

```

XX SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;
    Query Match 100.0%; Score 20; DB 22; Length 27;
    Best Local Similarity 100.0%; Pred. No. 0.5;
    Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCGTCGACGTCAGAGAG 20
Db 4 GATTGCGTCGACGTCAGAGAG 23

RESULT 11
AAH77397/c
ID AAH77397 standard; DNA; 27 BP.
XX AC AAH77397;
XX DT 05-NOV-2001 (first entry)
XX DE Cyclic AMP response element CRE consensus oligonucleotide probe #2.
XX KW Cancer; human; dithiocarbamate derivative; anticancer agent; probe; ss.
XX OS Unidentified.
XX PN US2001016600-A1.
XX PD 23-AUG-2001.
XX PF 12-DEC-2000; 2000US-0735205.
XX PR 08-SEP-1998; 98US-0099390.
XX PR 08-SEP-1999; 99US-0392122.
XX PR 05-OCT-2000; 2000US-0679932.
XX PA (KENN/) KENNEDY T P.
XX PI Kennedy TP;
XX PI WPI; 2001-557127/62.
XX DR Treating cancer, asthma and cancer and reducing hypoxic or ischemic
PT damage comprises administering dithiocarbamate thiolate anion or
PT dithiocarbamate thiolate metal complex.
XX PS Disclosure; Page 10; 36pp; English.
XX CC The present invention describes a method of treating cancer, asthma and
CC arthritis and reducing hypoxic or ischaemic damage, involving
CC administering a dithiocarbamate thiolate anion or metal ion complex to
CC the patient. The present sequence is a probe for the cyclic AMP response
CC element CRE, which was described in the exemplification of the invention.
XX SQ Sequence 27 BP; 5 A; 9 C; 5 G; 8 T; 0 other;
    Query Match 100.0%; Score 20; DB 22; Length 27;
    Best Local Similarity 100.0%; Pred. No. 0.5;
    Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCGTCGACGTCAGAGAG 20
Db 24 GATTGCGTCGACGTCAGAGAG 5

RESULT 12
AAH87956
ID AAH87956 standard; DNA; 27 BP.
XX AC AAH87956;
XX DT 20-JUL-2001 (first entry)
XX DE Cyclic AMP responsive element CRE consensus oligo for EMSA #2.
XX XX

DE XX Cyclic-AMP responsive element CRE consensus oligo for EMSA #1.
KW XX Cyclic-AMP responsive element; electrophoretic mobility shift assay;
KW EMSA; CRE; NF-kappaB; cancer; tetraethyl thiuram disulphide;
KW dithiocarbamate; tumour; metal ion; copper ion; cytokine;
KW ceruloplasmin; anticancer; cytostatic; ss.
XX OS Synthetic.
XX PN WO200117522-A1.
XX PD 15-MAR-2001.
XX PF 15-NOV-1999; 99WO-US27193.
XX PR 08-SEP-1999; 99US-0392122.
XX PA (CHAR-) CHARLOTTE-MECKLENBURG HOSPITAL AUTHORITY.
XX PI Kennedy TP;
XX DR WPI; 2001-281426/29.
XX PT Treating cancer, e.g. melanoma, lung cancer, breast cancer or prostatic
PT carcinoma, comprises administration of a thiuram disulfide optionally
PT in combination with a heavy metal ion, ceruloplasmin or a cytokine e.g.
PT interferon-alpha.
XX PS Disclosure; Page 24; 60pp; English.
XX CC The present invention describes a method for treating established cancer
CC in a mammal. The method comprises administering a thiuram disulfide (I).
CC (I) has anticancer and cytostatic activities. (I) induces apoptosis and
CC G2 cell-cycle arrest, by reducing expression of cyclin A (by preventing
CC binding of transcription factors to DNA regulatory elements involved in
CC control of cyclin A expression). The method can be used to treat cancers,
CC e.g. melanoma, non-small cell lung cancer, small cell lung cancer, renal
CC cancer, colorectal cancer, breast cancer, pancreatic cancer, gastric
CC cancer, bladder cancer, ovarian cancer, uterine cancer, lymphoma and
CC prostate cancer, especially melanoma, lung cancer, breast cancer and
CC prostate carcinoma. The tumour-inhibiting effect of (I) is dependent on
CC heavy metal ions, so administering (I) together with such ions (or with
CC their intracellular carriers, e.g. ceruloplasmin or with serum
CC ceruloplasmin stimulants, e.g. interferon-alpha) will increase the
CC antiproliferative/antineoplastic effect. (I) also potentiates the
CC effect of standard anticancer agents. (I) is already known for treating
CC alcohol abuse (inhibitor of aldehyde dehydrogenase) and is relatively
CC nontoxic and safe. The present sequence represents a cyclic-AMP
CC responsive element CRE consensus oligonucleotide for use in an
CC electrophoretic mobility shift assay (EMSA), which is used in the
CC exemplification of the present invention.
XX SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;
    Query Match 100.0%; Score 20; DB 22; Length 27;
    Best Local Similarity 100.0%; Pred. No. 0.5;
    Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GATTGCGTCGACGTCAGAGAG 20
Db 4 GATTGCGTCGACGTCAGAGAG 23

RESULT 13
AAH87957/c
ID AAH87957 standard; DNA; 27 BP.
XX AC AAH87957;
XX DT 20-JUL-2001 (first entry)
XX DE Cyclic-AMP responsive element CRE consensus oligo for EMSA #2.
XX XX

```

KW Cyclic-AMP responsive element; electrophoretic mobility shift assay;  
 KW EMSA; CRE; NF-kappaB; cancer; tetraethyl thiuram disulfide;  
 KW dithiocarbamate; tumor; metal ion; copper ion; cytokine;  
 KW ceruloplasmin; anticancer; cytostatic; ss.  
 XX Synthetic.  
 OS  
 XX WO200117522-A1.  
 PN  
 XX 15-MAR-2001.  
 PD  
 XX  
 PF 15-NOV-1999; 99WO-US27193.  
 XX  
 PR 08-SEP-1999; 99US-0392122.  
 XX  
 XX (CHAR-) CHARLOTTE-MECKLENBURG HOSPITAL AUTHORITY.  
 PA  
 XX Kennedy TP;  
 PI  
 XX WPI; 2001-281426/29.  
 DR  
 XX Treating cancer, e.g. melanoma, lung cancer, breast cancer or prostatic  
 PT carcinoma, comprises administration of a thiuram disulfide optionally  
 PT in combination with a heavy metal ion, ceruloplasmin or a cytokine e.g.  
 PT interferon-alpha.  
 PT  
 XX Disclosure; Page 24; 60pp; English.  
 PS  
 XX The present invention describes a method for treating established cancer  
 CC in a mammal. The method comprises administering a thiuram disulfide (I).  
 CC (I) has anticancer and cytostatic activities. (I) induces apoptosis and  
 CC G2 cell-cycle arrest, by reducing expression of cyclin A (by preventing  
 CC binding of transcription factors to DNA regulatory elements involved in  
 CC control of cyclin A expression). The method can be used to treat cancers,  
 CC e.g. melanoma, non-small cell lung cancer, small cell lung cancer, renal  
 CC cancer, colorectal cancer, breast cancer, pancreatic cancer, gastric  
 CC cancer, bladder cancer, ovarian cancer, uterine cancer, lymphoma and  
 CC prostate cancer, especially melanoma, lung cancer, breast cancer and  
 CC prostate carcinoma. The tumour-inhibiting effect of (I) is dependent on  
 CC heavy metal ions, so administering (I) together with such ions (or with  
 CC their intracellular carriers, e.g. ceruloplasmin or with serum  
 CC ceruloplasmin stimulants, e.g. interferon-alpha) will increase the  
 CC antiproliferative/antineoplastic affect. (I) also potentiates the  
 CC effect of standard anticancer agents. (I) is already known for treating  
 CC alcohol abuse (inhibitor of aldehyde dehydrogenase) and is relatively  
 CC nontoxic and safe. The present sequence represents a cyclic-AMP  
 CC responsive element CRE consensus oligonucleotide for use in an  
 CC electrophoretic mobility shift assay (EMSA), which is used in the  
 CC exemplification of the present invention.  
 XX  
 SQ Sequence 27 BP; 5 A; 9 C; 5 G; 8 T; 0 other;  
 Query Match 100.0%; Score 20; DB 22; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.5;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GATTGCTGACGTCAGAG 20  
 |||||  
 DB 24 GATTGCTGACGTCAGAG 5  
 RESULT 14  
 AAF76267  
 ID AAF76267 standard; DNA; 27 BP.  
 XX  
 AC AAF76267;  
 XX  
 DT 05-JUN-2001 (first entry)  
 XX  
 DE CAMP response element (CRE) competitor EMSA probe.  
 XX  
 KW NF-kappa-B; nuclear factor-kappa-B; CAMP response element; CRE;  
 KW nuclear translocation inhibition; heparin; internalisation;

KW NF-kappa-B dependent gene expression inhibition; cytokine;  
 KW tumour necrosis factor; TNF; interleukin; IL-1; IL-2; IL-6; IL-8;  
 KW interferon-beta; interferon-gamma; tissue factor-1; complement;  
 KW inducible nitric oxide synthase; diabetic vascular disease;  
 KW heart failure; asthma; sepsis; ischaemic-reperfusion injury;  
 KW electrophoretic mobility shift assay; competitor EMSA probe; ds.  
 XX Unidentified.  
 OS  
 XX WO200119376-A2.  
 PN  
 XX 22-MAR-2001.  
 PD  
 XX  
 PF 12-SEP-2000; 2000WO-US24910.  
 XX  
 PR 13-SEP-1999; 99US-0395081.  
 XX  
 XX (CHAR-) CHARLOTTE-MECKLENBURG HOSPITAL AUTHORITY.  
 PA  
 XX Kennedy TP;  
 PI  
 XX WPI; 2001-244698/25.  
 DR  
 XX Inhibiting NF-kappa-B activity, useful for treating e.g. diabetic  
 XX vascular disease, heart failure, asthma and sepsis, comprises  
 PT administering heparin to cells in patient to inhibit translocation of  
 PT NF-kappa-B from cytoplasm to nucleus.  
 PT  
 XX Examples; Page 22; 68pp; English.  
 PS  
 XX The invention relates to a method of inhibiting nuclear factor-kappa-B  
 CC (NF-kappa-B) activity in a patient, comprising the administration of  
 CC heparin to the cells in the patient, such that the heparin is  
 CC internalised into the cytoplasm of cells in the patient. The invention  
 CC is based on the discovery that heparin is able to block the  
 CC translocation of NF-kappa-B from the cytoplasm to the nucleus. This in  
 CC turn inhibits NF-kappa-B dependent gene expression. Such NF-kappa-B  
 CC dependent genes include genes encoding cytokines such as tumour necrosis  
 CC factor (TNF), IL-1 (interleukin-1), IL-2, IL-6, IL-8, interferon-beta,  
 CC interferon-gamma, tissue factor-1, complement and inducible nitric  
 CC oxide synthase. The method of the invention is used for treating or  
 CC preventing diabetic vascular disease, heart failure, asthma, sepsis and  
 CC ischaemic-reperfusion injury. Heparin may be administered in combination  
 CC with other active agents that treat or prevent another disease or  
 CC symptom in the patient, e.g., antiviral agents, antibiotics, antifungal  
 CC agents and antiinflammatory agents. The method of the invention offers  
 CC significant advantages over prior art treatments for the above  
 CC conditions. Heparin is relatively non-toxic and safe, and should not  
 CC produce the side effects such as hypertension, glucose intolerance  
 CC and bone demineralisation that are encountered with the use of  
 CC glucocorticoids for blocking the NF-kappa-B nuclear translocation.  
 CC Additionally, heparin is readily available and easily used. Sequences  
 CC AAF76266-AAF76267 represents EMSA (electrophoretic mobility shift assay)  
 CC probes used to measure the effect of heparin on NF-kappa-B nuclear  
 CC translocation. EMSA probe AAF76266 comprises a consensus NF-kappa-B  
 CC response element, and EMSA competitor probe AAF76267 comprises a  
 CC CAMP response element (CRE).  
 XX  
 SQ Sequence 27 BP; 8 A; 5 C; 9 G; 5 T; 0 other;

Query Match 100.0%; Score 20; DB 22; Length 27;  
 Best Local Similarity 100.0%; Pred. No. 0.5;  
 Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
 QY 1 GATTGCTGACGTCAGAG 20  
 |||||  
 DB 4 GATTGCTGACGTCAGAG 23  
 RESULT 15  
 ABA92274  
 ID ABA92274 standard; DNA; 27 BP.  
 XX





us-09-355-254f-17.rni

Thu Dec 12 07:53:51 2002

GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model  
Run on: December 12, 2002, 01:05:40 : Search time 22.2464 Seconds  
(without alignments)  
275.709 Million cell updates/sec.

Title: US-09-355-254F-17

Perfect score: 20  
Sequence: 1 tatgcataattcctgtaagt 20

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 441362 seqs, 1533381 residues 687286  
Total number of hits satisfying chosen parameters:

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

- Database :
- 1: /cgn2\_6/ptodata/1/ina/5A\_COMB.seq.\*
  - 2: /cgn2\_6/ptodata/1/ina/5B\_COMB.seq.\*
  - 3: /cgn2\_6/ptodata/1/ina/6A\_COMB.seq.\*
  - 4: /cgn2\_6/ptodata/1/ina/6B\_COMB.seq.\*
  - 5: /cgn2\_6/ptodata/1/ina/PCTUS\_COMB.seq.\*
  - 6: /cgn2\_6/ptodata/1/ina/backfiles1.seq.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	16	80.0	19	1	US-08-410-780A-75
2	16	80.0	19	1	US-08-411-020-40
3	16	80.0	19	5	PCT-US95-04511-75
4	15	75.0	17	1	US-08-369-796-21
5	15	75.0	17	2	US-08-852-091-21
6	15	75.0	17	5	PCT-US95-17025-21
7	15	75.0	19	1	US-08-410-780A-75
8	15	75.0	19	1	US-08-411-020-41
9	15	75.0	19	5	PCT-US95-04511-76
10	14.8	74.0	47	1	US-08-171-389-37
11	14.8	74.0	47	2	US-08-123-936-37
12	14.8	74.0	47	3	US-08-475-228A-37
13	14.8	74.0	47	3	US-08-482-080A-37
14	14.8	74.0	47	4	US-09-354-947-37
15	14.8	74.0	47	5	PCT-US93-12388-37
16	14.8	74.0	50	1	US-08-171-389-437
17	14.8	74.0	50	1	US-08-123-936-437
18	14.8	74.0	50	1	US-08-475-228A-437
19	14.8	74.0	50	2	US-08-475-228A-437
20	14.8	74.0	50	2	US-08-482-080A-437
21	14.8	74.0	50	3	US-08-482-080A-437
22	14.8	74.0	50	3	US-08-482-080A-437
23	14.8	74.0	50	4	US-09-354-947-437
24	14.8	74.0	50	4	US-09-354-947-437
25	14.8	74.0	50	5	PCT-US93-12388-437
26	14.8	74.0	50	5	PCT-US93-12388-437
27	14.8	74.0	50	5	PCT-US93-12388-437

28	14.4	72.0	19	1	US-08-410-780A-23	Sequence 23, Appl
29	14.4	72.0	19	1	US-08-410-780A-25	Sequence 25, Appl
30	14.4	72.0	19	1	US-08-410-780A-33	Sequence 33, Appl
31	14.4	72.0	19	1	US-08-410-780A-49	Sequence 49, Appl
32	14.4	72.0	19	1	US-08-410-780A-51	Sequence 51, Appl
33	14.4	72.0	19	1	US-08-410-780A-55	Sequence 55, Appl
34	14.4	72.0	19	1	US-08-411-020-42	Sequence 42, Appl
35	14.4	72.0	19	5	PCT-US95-04511-23	Sequence 23, Appl
36	14.4	72.0	19	5	PCT-US95-04511-25	Sequence 25, Appl
37	14.4	72.0	19	5	PCT-US95-04511-33	Sequence 33, Appl
38	14.4	72.0	19	5	PCT-US95-04511-49	Sequence 49, Appl
39	14.4	72.0	19	5	PCT-US95-04511-51	Sequence 51, Appl
40	14.4	72.0	19	5	PCT-US95-04511-55	Sequence 55, Appl
41	14	70.0	20	4	US-09-489-868A-31	Sequence 31, Appl
42	13.4	67.0	19	1	US-08-410-780A-24	Sequence 24, Appl
43	13.4	67.0	19	1	US-08-410-780A-26	Sequence 26, Appl
44	13.4	67.0	19	1	US-08-410-780A-34	Sequence 34, Appl
45	13.4	67.0	19	1	US-08-410-780A-50	Sequence 50, Appl

ALIGNMENTS

RESULT 1  
US-08-410-780A-75 : Application US/08410780A  
; Patent No. 5707803  
; GENERAL INFORMATION:  
; APPLICANT: LAMB, I. PETER  
; APPLICANT: SEIDEL, H. MARTI  
; TITLE OF INVENTION: DNA REGULATORY ELEMENTS RESPONSIVE TO  
; TITLE OF INVENTION: CYTOKINES AND METHODS FOR THEIR USE  
; NUMBER OF SEQUENCES: 77  
; CORRESPONDENCE ADDRESS:  
; ADDRESSEE: LIGAND PHARMACEUTICALS INCORPORATED  
; STREET: 9393 TOWNE CENTRE DRIVE  
; CITY: SAN DIEGO  
; STATE: CALIFORNIA  
; COUNTRY: US  
; ZIP: 92121

COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.30  
CURRENT APPLICATION NUMBER: US/08/410,780A  
FILING DATE: 27-MAR-1995  
CLASSIFICATION: 435

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/228,934  
FILING DATE: 14-APR-1994  
ATTORNEY/AGENT INFORMATION:  
NAME: JURGENSEN, THOMAS E  
REGISTRATION NUMBER: 34,195  
REFERENCE/DOCKET NUMBER: 016-0017A.US  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (619) 550-7675  
TELEFAX: (619) 535-3906  
INFORMATION FOR SEQ ID NO: 75:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 19 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: single  
TOPOLOGY: linear

MOLECULE TYPE: other nucleic acid  
DESCRIPTION: /desc = "OTHER NUCLEIC ACID,  
SYNTHETIC DNA"  
US-08-410-780A-75

Query Match 80.0%; Score 16; DB 1; Length 19;  
Best Local Similarity 100.0%; Pred No. 15;  
Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;



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Query Match 75.0%; Score 15; DB 1; Length 17;
Best Local Similarity 100.0%; Pred. No. 47;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
DB 1 ATATTCCTGTAAGTG 15

RESULT 5
US-08-852-091-21
; Sequence 21, Application US/08852091
; Patent No. 5863228
; GENERAL INFORMATION:
; APPLICANT: James E. Darnell, Jr.
; APPLICANT: Zilong Wen
; APPLICANT: Curt M. Horvath
; APPLICANT: Zhong Zhong
; TITLE OF INVENTION: FUNCTIONALLY ACTIVE REGIONS OF SIGNAL
; TITLE OF INVENTION: TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION (STAT) PROTEINS
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Klauber & Jackson
; STREET: 411 Hackensack Avenue
; CITY: Hackensack
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07601
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/852,091
; FILING DATE: 06-MAY-1997
; CLASSIFICATION: 424
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/369,796
; FILING DATE: 06-JAN-1995
; ATTORNEY/AGENT INFORMATION:
; NAME: Jackson Esq., David A.
; REGISTRATION NUMBER: 26,742
; REFERENCE/DOCKET NUMBER: 600-1-116
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 201 487-5800
; TELEFAX: 201 343-1684
; TELEX: 133521
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA synthetic probe
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; US-08-852-091-21

Query Match 75.0%; Score 15; DB 2; Length 17;
Best Local Similarity 100.0%; Pred. No. 47;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
DB 1 ATATTCCTGTAAGTG 15

RESULT 6
PCT-US95-17025-21
; Sequence 21, Application PC/TUS9517025
; GENERAL INFORMATION:
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; APPLICANT: James E. Darnell, Jr.
; APPLICANT: Zilong Wen
; APPLICANT: Curt M. Horvath
; APPLICANT: Zhong Zhong
; TITLE OF INVENTION: FUNCTIONALLY ACTIVE REGIONS OF SIGNAL
; TITLE OF INVENTION: TRANSDUCER AND ACTIVATOR OF TRANSCRIPTION (STAT) PROTEINS
; NUMBER OF SEQUENCES: 39
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Klauber & Jackson
; STREET: 411 Hackensack Avenue
; CITY: Hackensack
; STATE: New Jersey
; COUNTRY: USA
; ZIP: 07601
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: PCT/US95/17025
; FILING DATE: 28-DEC-1995
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/369,796
; FILING DATE: 06-JAN-1995
; CLASSIFICATION:
; ATTORNEY/AGENT INFORMATION:
; NAME: Jackson Esq., David A.
; REGISTRATION NUMBER: 26,742
; REFERENCE/DOCKET NUMBER: 600-1-116
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 201 487-5800
; TELEFAX: 201 343-1684
; TELEX: 133521
; INFORMATION FOR SEQ ID NO: 21:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 17 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: DNA synthetic probe
; HYPOTHETICAL: NO
; ANTI-SENSE: NO
; PCT-US95-17025-21

Query Match 75.0%; Score 15; DB 5; Length 17;
Best Local Similarity 100.0%; Pred. No. 47;
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATTCCTGTAAGTG 20
DB 1 ATATTCCTGTAAGTG 15

RESULT 7
US-08-410-780A-76/c
; Sequence 76, Application US/08410780A
; Patent No. 5707803
; GENERAL INFORMATION:
; APPLICANT: LAMB, I. PETER
; APPLICANT: SEIDEL, H. MARTI
; TITLE OF INVENTION: DNA REGULATORY ELEMENTS RESPONSIVE TO
; TITLE OF INVENTION: CYTOKINES AND METHODS FOR THEIR USE
; NUMBER OF SEQUENCES: 77
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: LIGAND PHARMACEUTICALS INCORPORATED
; STREET: 9393 TOWNE CENTRE DRIVE
; CITY: SAN DIEGO
; STATE: CALIFORNIA
; COUNTRY: US
; ZIP: 92121
; COMPUTER READABLE FORM:
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;; MEDIUM TYPE: Floppy disk  
;; COMPUTER: IBM PC compatible  
;; OPERATING SYSTEM: PC-DOS/MS-DOS  
;; SOFTWARE: PatentIn Release #1.0, Version #1.30  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: US/08/410,780A  
;; FILING DATE: 27-MAR-1995  
;; CLASSIFICATION: 435  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 08/228,934  
;; FILING DATE: 14-APR-1994  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: JURGENSEN, THOMAS E  
;; REGISTRATION NUMBER: 34,195  
;; REFERENCE/DOCKET NUMBER: 016-0017A.US  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: (619) 550-7675  
;; TELEFAX: (619) 535-3906  
;; INFORMATION FOR SEQ ID NO: 76:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 19 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: single  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: other nucleic acid  
;; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,  
;; SYNTHETIC DNA"  
US-08-410-780A-76  
  
Query Match 75.0%; Score 15; DB 1; Length 19;  
Best Local Similarity 100.0%; Pred. No. 48;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 6 ATATTCCTGTAAGTG 20  
Db 19 ATATTCCTGTAAGTG 5  
  
RESULT 8  
US-08-411-020-41/c  
;; Sequence 41, Application US/08411020  
;; Patent No. 5712094  
;; GENERAL INFORMATION:  
;; APPLICANT: SEIDEL, H. MARTI  
;; APPLICANT: LAMB, I. PETER  
;; APPLICANT: CHAN, SHIN-SHAY TIAN  
;; TITLE OF INVENTION: METHODS AND ASSOCIATED REAGENTS FOR  
;; DETECTING MODULATORS OF CYTOKINE ACTION  
;; NUMBER OF SEQUENCES: 59  
;; CORRESPONDENCE ADDRESS:  
;; ADDRESSEE: Ligand Pharmaceuticals Incorporated  
;; STREET: 9393 Towne Centre Drive  
;; CITY: San Diego  
;; STATE: California  
;; COUNTRY: US  
;; ZIP: 92121  
;; MEDIUM TYPE: Floppy disk  
;; COMPUTER: IBM PC compatible  
;; OPERATING SYSTEM: PC-DOS/MS-DOS  
;; SOFTWARE: PatentIn Release #1.0, Version #1.30  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: US/08/411,020  
;; FILING DATE: 27-MAR-1995  
;; CLASSIFICATION: 435  
;; ATTORNEY/AGENT INFORMATION:  
;; NAME: Jurgensen, Thomas E.  
;; REGISTRATION NUMBER: 34,195  
;; REFERENCE/DOCKET NUMBER: 016-0030.US  
;; TELECOMMUNICATION INFORMATION:  
;; TELEPHONE: (619) 550-7675  
;; TELEFAX: (619) 535-3906  
;; INFORMATION FOR SEQ ID NO: 41:

;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 19 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: single  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: other nucleic acid  
;; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,  
;; SYNTHETIC DNA"  
US-08-411-020-41  
  
Query Match 75.0%; Score 15; DB 1; Length 19;  
Best Local Similarity 100.0%; Pred. No. 48;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 6 ATATTCCTGTAAGTG 20  
Db 19 ATATTCCTGTAAGTG 5  
  
RESULT 9  
PCT-US95-04511-76/c  
;; Sequence 76, Application PC/TUS9504511  
;; GENERAL INFORMATION:  
;; APPLICANT:  
;; TITLE OF INVENTION: DNA REGULATORY ELEMENTS RESPONSIVE TO  
;; CYTOKINES AND METHODS FOR THEIR USE  
;; NUMBER OF SEQUENCES: 76  
;; COMPUTER READABLE FORM:  
;; MEDIUM TYPE: Floppy disk  
;; COMPUTER: IBM PC compatible  
;; OPERATING SYSTEM: PC-DOS/MS-DOS  
;; SOFTWARE: PatentIn Release #1.0, Version #1.30 (EPO)  
;; CURRENT APPLICATION DATA:  
;; APPLICATION NUMBER: PCT/US95/04511  
;; FILING DATE:  
;; CLASSIFICATION:  
;; PRIOR APPLICATION DATA:  
;; APPLICATION NUMBER: US 08/228,934  
;; FILING DATE: 14-APR-1994  
;; INFORMATION FOR SEQ ID NO: 76:  
;; SEQUENCE CHARACTERISTICS:  
;; LENGTH: 19 base pairs  
;; TYPE: nucleic acid  
;; STRANDEDNESS: single  
;; TOPOLOGY: linear  
;; MOLECULE TYPE: other nucleic acid  
;; DESCRIPTION: /desc = "OTHER NUCLEIC ACID,  
;; SYNTHETIC DNA"  
PCT-US95-04511-76  
  
Query Match 75.0%; Score 15; DB 5; Length 19;  
Best Local Similarity 100.0%; Pred. No. 48;  
Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
  
QY 6 ATATTCCTGTAAGTG 20  
Db 19 ATATTCCTGTAAGTG 5  
  
RESULT 10  
US-08-171-389-37/c  
;; Sequence 37, Application US/08171389  
;; Patent No. 5578444  
;; GENERAL INFORMATION:  
;; APPLICANT: Edwards, Cynthia A.  
;; APPLICANT: Cantor, Charles R.  
;; APPLICANT: Andrews, Beth M.  
;; APPLICANT: Turin, Lisa M.  
;; APPLICANT: Fry, Kirk E.  
;; TITLE OF INVENTION: Sequence-Directed DNA Binding  
;; MOLECULES, COMPOSITIONS AND METHODS  
;; NUMBER OF SEQUENCES: 641  
;; CORRESPONDENCE ADDRESS:

ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/171,389  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/081,070  
FILING DATE: 22-JUN-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Fabian, Gary R.  
REGISTRATION NUMBER: 33,875  
REFERENCE/DOCKET NUMBER: 4600-0175/G19P3  
TELEPHONE: (415) 324-0880  
TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 37:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Human alpha-amylase gene  
US-08-171-389-37

Query Match 74.0%; Score 14.8; DB 1; Length 47;  
Best Local Similarity 88.9%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
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Db 28 TATTATATTCCTGTAAG 11

RESULT 11  
US-08-123-936-37/c  
Sequence 37, Application US/08123936  
Patent No. 5726014  
GENERAL INFORMATION:  
APPLICANT: Edwards, Cynthia A.  
APPLICANT: Cantor, Charles R.  
APPLICANT: Andrews, Beth M.  
APPLICANT: Turin, Lisa M.  
TITLE OF INVENTION: Screening Assay for the Detection of  
NUMBER OF SEQUENCES: 640  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA

ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/123,936  
FILING DATE:  
CLASSIFICATION: 435  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
ATTORNEY/AGENT INFORMATION:  
NAME: Fabian, Gary R.  
REGISTRATION NUMBER: 33,875  
REFERENCE/DOCKET NUMBER: 4600-0075.32/G19P2  
TELEPHONE: (415) 324-0880  
TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 37:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Human alpha-amylase gene  
US-08-123-936-37

Query Match 74.0%; Score 14.8; DB 1; Length 47;  
Best Local Similarity 88.9%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
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Db 28 TATTATATTCCTGTAAG 11

RESULT 12  
US-08-475-228A-37/c  
Sequence 37, Application US/08475228A  
Patent No. 5869241  
GENERAL INFORMATION:  
APPLICANT: Edwards, Cynthia A.  
APPLICANT: Cantor, Charles R.  
APPLICANT: Andrews, Beth M.  
APPLICANT: Turin, Lisa M.  
APPLICANT: Fry, Kirk E.  
TITLE OF INVENTION: Sequence-Directed DNA Binding  
NUMBER OF SEQUENCES: 664  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/475,228A  
FILING DATE: 06-JUN-1995  
PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/081,070  
FILING DATE: 22-JUN-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Stratford, Carol A.  
REGISTRATION NUMBER: 34,444  
REFERENCE/DOCKET NUMBER: 4600-0175.21/G19P3D2  
TELEPHONE: (415) 324-0880  
TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 37:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Human alpha-amylase gene  
US-08-475-228A-37

Query Match 74.0%; Score 14.8; DB 2; Length 47;  
Best Local Similarity 88.9%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
||| |||||  
Db 28 TATTATATTCCTGTAAG 11

RESULT 13  
US-08-482-080A-37/C  
Sequence 37, Application US/08482080A  
Patent No. 6010849  
GENERAL INFORMATION:  
APPLICANT: Edwards, Cynthia A.  
APPLICANT: Cantor, Charles R.  
APPLICANT: Andrews, Beth M.  
APPLICANT: Turin, Lisa M.  
APPLICANT: Fry, Kirk E.  
TITLE OF INVENTION: Sequence-Directed DNA Binding  
Molecules, Compositions and Methods  
NUMBER OF SEQUENCES: 664  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/08/482,080A  
FILING DATE: 07-JUN-1995  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/171,389  
FILING DATE: 20-DEC-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/081,070  
FILING DATE: 22-JUN-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Brady, John F.  
REGISTRATION NUMBER: 39,118  
REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1  
TELEPHONE: (650) 324-0880  
TELEFAX: (650) 324-0960  
INFORMATION FOR SEQ ID NO: 37:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Human alpha-amylase gene  
US-08-482-080A-37

Query Match 74.0%; Score 14.8; DB 3; Length 47;  
Best Local Similarity 88.9%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
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Db 28 TATTATATTCCTGTAAG 11

RESULT 14  
US-09-354-947-37/C  
Sequence 37, Application US/09354947  
Patent No. 6384208  
GENERAL INFORMATION:  
APPLICANT: Edwards, Cynthia A.  
APPLICANT: Cantor, Charles R.  
APPLICANT: Andrews, Beth M.  
APPLICANT: Turin, Lisa M.  
APPLICANT: Fry, Kirk E.  
TITLE OF INVENTION: Sequence-Directed DNA Binding  
Molecules, Compositions and Methods  
NUMBER OF SEQUENCES: 664  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: US/09/354,947  
FILING DATE:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/482,080  
FILING DATE: 07-JUN-1995  
APPLICATION NUMBER: US 08/171,389  
FILING DATE: 20-DEC-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993

PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/723,618  
FILING DATE: 27-JUN-1991  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/081,070  
FILING DATE: 22-JUN-1993  
ATTORNEY/AGENT INFORMATION:  
NAME: Brady, John F.  
REGISTRATION NUMBER: 39,118  
REFERENCE/DOCKET NUMBER: 4600-0175.20/G19P3D1  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (650) 324-0880  
TELEFAX: (650) 324-0960  
INFORMATION FOR SEQ ID NO: 37:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Human alpha-amylase gene  
US-09-354-947-37

Query Match 74.0%; Score 14.8; DB 4; Length 47;  
Best Local Similarity 88.9%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
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Db 28 TATTATATTCCTGTAAG 11

RESULT 15  
PCT-US93-12388-37/c  
Sequence 37, Application PC/TUS9312388  
GENERAL INFORMATION:  
APPLICANT:  
TITLE OF INVENTION: Sequence-Directed DNA Binding  
TITLE OF INVENTION: Molecules, Compositions and Methods  
NUMBER OF SEQUENCES: 641  
CORRESPONDENCE ADDRESS:  
ADDRESSEE: Genelabs Technologies, Inc.  
STREET: 505 Penobscot Drive  
CITY: Redwood City  
STATE: CA  
COUNTRY: USA  
ZIP: 94063  
COMPUTER READABLE FORM:  
MEDIUM TYPE: Floppy disk  
COMPUTER: IBM PC compatible  
OPERATING SYSTEM: PC-DOS/MS-DOS  
SOFTWARE: PatentIn Release #1.0, Version #1.25  
CURRENT APPLICATION DATA:  
APPLICATION NUMBER: PCT/US93/12388  
FILING DATE:  
CLASSIFICATION:  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 08/123,936  
FILING DATE: 17-SEP-1993  
PRIOR APPLICATION DATA:  
APPLICATION NUMBER: US 07/996,783  
FILING DATE: 23-DEC-1992  
ATTORNEY/AGENT INFORMATION:  
NAME: Fabian, Gary R.  
REGISTRATION NUMBER: 33,875  
REFERENCE/DOCKET NUMBER: 4600-0175.41/G19PCT2  
TELECOMMUNICATION INFORMATION:  
TELEPHONE: (415) 324-0880

TELEFAX: (415) 324-0960  
INFORMATION FOR SEQ ID NO: 37:  
SEQUENCE CHARACTERISTICS:  
LENGTH: 47 base pairs  
TYPE: nucleic acid  
STRANDEDNESS: double  
TOPOLOGY: linear  
MOLECULE TYPE: DNA (genomic)  
HYPOTHETICAL: NO  
ORIGINAL SOURCE:  
INDIVIDUAL ISOLATE: Human alpha-amylase gene  
PCT-US93-12388-37  
Query Match 74.0%; Score 14.8; DB 5; Length 47;  
Best Local Similarity 88.9%; Pred. No. 67;  
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
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Db 28 TATTATATTCCTGTAAG 11

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Job time : 22.2464 secs

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GenCore version 5.1.1.3  
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OM nucleic - nucleic search, using sw model

Run on: December 12, 2002, 01:04:50 ; Search time 102.319 seconds  
(without alignments)  
440.192 Million cell updates/sec

Title: US-09-355-254F-17

Perfect score: 20  
Sequence: 1 tatgcataattctgtaagt 20

Scoring table: IDENTITY\_NUC  
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Searched: 2185239 seqs, 112599159 residues

Total number of hits satisfying chosen parameters: 2390332

Minimum DB seq length: 0  
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Maximum Match 100%  
Listing first 45 summaries

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- 24: /SIDS2/gcgdata/geneseq/geneseq-emb1/NA2002.DAT.\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
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2	20	100.0	20	24	AA139193
3	19	95.0	24	24	ABN80818
4	16	80.0	19	17	AA141604
5	15	75.0	17	17	AA131283
6	14.8	74.0	47	15	AAQ69287
7	14.8	74.0	47	18	AA163749
8	14.8	74.0	47	20	AA17037
9	14.8	74.0	47	24	ABR82528

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C 11	14.8	74.0	50	15	AAQ69687	Human pancreatic a
C 12	14.8	74.0	50	18	AA164148	Human pancreatic a
C 13	14.8	74.0	50	18	AA164149	Human pancreatic a
C 14	14.8	74.0	50	20	AA117436	Test sequence from
C 15	14.8	74.0	50	20	AA117437	DNA binding molecu
C 16	14.8	74.0	50	24	ABK82927	DNA binding molecu
C 17	14.8	74.0	50	24	ABK82928	Oligonucleotide co
C 18	14.4	72.0	19	17	AA141605	Human cot oncogene
C 19	14	70.0	20	22	AA111323	Primer 2 for pMAL-
C 20	14	70.0	27	15	AA114907	C2P-3 Gene primer.
C 21	14	70.0	27	15	AAQ55426	Single nucleotide
C 22	13.8	69.0	22	21	AA171335	Humanised anti-Fas
C 23	13.6	68.0	50	19	AAV66651	Humanised anti-Fas
C 24	13.6	68.0	50	21	AA178342	PEBP2 alpha A gene
C 25	13.4	67.0	21	20	AA133318	Oligonucleotide ad
C 26	13.4	67.0	24	24	ABQ01109	Oligonucleotide ad
C 27	13.4	67.0	24	24	ABQ06151	Oligonucleotide ad
C 28	13.4	67.0	24	24	ABQ06192	SNP specific lower
C 29	13.4	67.0	25	22	AA139126	Human silent SNP c
C 30	13.4	67.0	51	22	AA179532	5' fragment of the
C 31	13.4	67.0	60	14	AAQ51030	Human spliced tran
C 32	13.4	67.0	60	24	ABN39544	Sequencing primer
C 33	13.2	66.0	21	22	AA187028	Human genomic DNA
C 34	13.2	66.0	31	20	AA138743	Human spliced tran
C 35	13.2	66.0	60	24	ABN36070	Rat spliced trans
C 36	13.2	66.0	65	24	ABN29969	Human 5' EST isola
C 37	13.2	66.0	99	21	AA242637	Ly6 GAS regulatory
C 38	13	65.0	13	16	AA102747	Human map-related
C 39	13	65.0	47	21	AA266670	Mouse spliced tran
C 40	13	65.0	65	24	ABN52711	Human genome biall
C 41	12.8	64.0	23	20	AA152734	PCR primer #2 for
C 42	12.8	64.0	36	13	AAQ31180	Citrate synthase 1
C 43	12.8	64.0	41	24	ABK48179	Human genome biall
C 44	12.8	64.0	47	20	AA152584	Human genome biall
C 45	12.8	64.0	47	20	AA152534	Human genome biall

## ALIGNMENTS

RESULT 1  
AAV46009  
ID AAV46009 standard; DNA; 20 BP.

XX AAV46009;

XX 16-OCT-1998 (first entry)

DT Immune adjuvant STAT1.

DE Immune system; adjuvant; vaccine; cancer; prophylactic; pathogenicity;

XX modulator; tolerance; regulator; helper cell; antigen; immunoglobulin;

KW Ig class; autoimmune response; T-cell; B-cell; tumour; ss.

KW Class Bacteria.

XX EP855184-A1.

XX 29-JUL-1998.

XX 23-JAN-1997; 97EP-0101019.

XX 23-JAN-1997; 97EP-0101019.

XX (HEG/) HEG K.

PA (LIPF/) LIPFORD G B.

PA (WAGN/) WAGNER H.

XX Heeg K, Lipford GB, Wagner H;

PI WPI; 1998-389630/34.

XX

PT Antigenic composition comprises polynucleotide fragment and antigen  
PT - used as vaccine to treat or prevent e.g. cancer or pathogen  
PT infections and to modulate immune response e.g. tolerance break and  
PT regulation of TH1/TH2 cells  
XX  
XX Example 5; Page 9; 28pp; English.  
PS  
XX AAV5993-V46019 are fragments of bacterial polynucleotides which are  
CC used as immune adjuvants for inclusion into vaccines to treat cancer and  
CC for prophylaxis and/or treatment of conditions caused by pathogenic  
CC micro-organisms. The polynucleotide is used for modulation of an immune  
CC response and the modulation is selected from the group break of  
CC tolerance, regulation of TH1/TH2 helper cell responses, switch of Ig  
CC classes, treatment of autoimmune responses and induction of tolerances.  
CC DNA oligomers are used to enhance the reactivity of immune cells to  
CC viral, bacterial and parasitic antigens, to break tolerance in anergic T  
CC and B cells e.g. against tumour antigens, as adjuvants in vaccination  
CC against tumour-defined antigens and immunostimulatory substances in an  
CC immune response against tumours and to suppress immune reactions of the  
CC innate and acquired immune system. The composition is inexpensive and  
CC stable and does not cause lethal shock, which happens with prior art  
CC bacterial sequences.  
XX  
SQ Sequence 20 BP; 5 A; 3 C; 4 G; 8 T; 0 other;  
Query Match 100.0%; Score 20; DB 19; Length 20;  
Best Local Similarity 100.0%; Pred. No. 1.3;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 TATGCATATTCCTGTAAGTG 20  
Db | | | | | | | | | | | | | | | | | | | |  
1 TATGCATATTCCTGTAAGTG 20  
RESULT 2  
AAL39193  
ID AAL39193 standard; DNA; 20 BP.  
XX  
AC AAL39193;  
XX  
DT 05-SEP-2002 (first entry)  
XX  
DE Murine Toll-like receptor related CpG DNA SEQ ID No 68.  
XX  
XX Murine Toll-like receptor; TLR9; TLR7; TLR8; ISNA; ds.  
KW  
XX Unidentified.  
OS  
XX WO200222809-A2.  
PN  
XX 21-MAR-2002.  
PD  
XX 17-SEP-2001; 2001WO-US29229.  
XX  
XX 15-SEP-2000; 2000US-233035P.  
PR  
XX 23-JAN-2001; 2001US-263657P.  
PR  
XX 17-MAY-2001; 2001US-291726P.  
PR  
XX 22-JUN-2001; 2001US-300210P.  
XX  
XX (COLE-) COLEY PHARM GMBH.  
XX  
XX Bauer S, Lipford G, Wagner H;  
PI  
XX WPI; 2002-393964/42.  
XX  
XX New isolated murine Toll-like receptor (TLR)9, TLR7, TLR8 polypeptides,  
PT useful for identifying species specificity of immunostimulatory nucleic  
PT acid and identifying immunostimulatory nucleic acids  
XX  
XX Disclosure: Page 76; 195pp; English.  
PS  
XX The invention relates to isolated murine Toll-like receptors (TLR)9,  
CC TLR7 and TLR8 polypeptides. These polypeptides comprise fully defined

CC sequences of 1032, 1050 or 1032 amino acids as given in specification, or  
CC their fragments, where TLR9, TLR7 and TLR8 polypeptides or their  
CC fragments have an amino acid sequence which is identical to human TLR9,  
CC TLR7 or TLR8 polypeptides or their fragment except for at least one amino  
CC acid of a murine TLR polypeptide. The isolated nucleic acids of the  
CC invention are useful for inhibiting TLR9 signalling activity in a cell.  
CC TLR7, TLR8 and TLR9 polypeptides are useful for identifying nucleic acid  
CC molecules which interact with a TLR polypeptide or its fragment. The  
CC TLR7, TLR8 and TLR9 polypeptides are also useful for identifying ISNA. The  
CC signalling activity of a test compound (that is not a nucleic acid, and  
CC is a polypeptide or a part of a combinatorial library of compounds) with  
CC an ISNA. The TLR7, TLR8 and TLR9 polypeptides are also useful for  
CC identifying species specificity of an ISNA. The isolated nucleic acids of  
CC the invention are useful as probes or primers. This polynucleotide  
CC sequence represents DNA relating to the isolated Toll-like receptors of  
CC the invention.  
XX  
SQ Sequence 20 BP; 5 A; 3 C; 4 G; 8 T; 0 other;  
Query Match 100.0%; Score 20; DB 24; Length 20;  
Best Local Similarity 100.0%; Pred. No. 1.3;  
Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;  
QY 1 TATGCATATTCCTGTAAGTG 20  
Db | | | | | | | | | | | | | | | | | | | |  
1 TATGCATATTCCTGTAAGTG 20  
RESULT 3  
ABN80818  
ID ABN80818 standard; DNA; 24 BP.  
XX  
AC ABN80818;  
XX  
DT 15-JUL-2002 (first entry)  
XX  
DE Human STAT-1 inhibitor oligonucleotide SEQ ID NO 36.  
XX  
XX Human; IRF-1; transcription factor; interferon regulatory factor;  
KW antisense; gene therapy; cardiovascular; transplant rejection;  
KW immunological hypersensitivity; asthma; inflammatory disease; psoriasis;  
KW Crohn's disease; autoimmune disease; diabetes mellitus;  
KW multiple sclerosis; rheumatoid arthritis; Th1 response; Th2 response;  
KW vasotropic; immunosuppressive; antiasthmatic; dermatological;  
KW anti-allergic; anti-ulcer; anti-inflammatory; antipsoriatic; antidiabetic;  
KW neuroprotective; antirheumatic; antiarthritic; ss.  
XX  
XX Homo sapiens.  
OS  
XX Synthetic.  
XX  
XX WO200229044-A2.  
PN  
XX 11-APR-2002.  
PD  
XX 04-OCT-2001; 2001WO-DE03835.  
XX  
XX 06-OCT-2000; 2000DE-1049549.  
PR  
XX 29-NOV-2000; 2000DE-1059144.  
PR  
XX (HECK/) HECKER M.  
PA (WAGN/) WAGNER A H.  
XX  
XX Hecker M, Wagner AH;  
PI  
XX WPI; 2002-383335/41.  
DR  
XX Inhibitor of the transcription factor IFR-1, useful for treating e.g.  
PT transplant rejection and autoimmune disease, reduces expression of CD40  
PT  
XX Example 3; Page 19; 45pp; German.  
PS  
XX

CC The invention relates to an inhibitor (I) of the expression and/or  
 CC activity of the transcription factor (IRF-1; interferon regulatory  
 CC factor) as a therapeutic agent, especially an oligonucleotide inhibitor  
 CC (ABN80783-ABN80804) or antisense oligonucleotide (ABN80805-ABN80808) used  
 CC in antisense gene therapy. (I) are used to prevent or treat  
 CC cardiovascular complications such as restenosis after angioplasty or  
 CC stenosis of venous by-passes, chronic or acute transplant rejection and  
 CC graft versus host disease, immunological hypersensitivity, e.g. bronchial  
 CC asthma or atopic dermatitis, inflammatory diseases such as ulcerative  
 CC colitis, psoriasis and Crohn's disease and autoimmune diseases such as  
 CC diabetes mellitus, multiple sclerosis, collagenosis (e.g. systemic lupus  
 CC erythematosus), rheumatoid arthritis and vasculitis. (I) simultaneously  
 CC weaken both Th1 and Th2 responses. The present sequence is that of an  
 CC oligonucleotide, useful to the invention.

XX  
 SQ Sequence 24 BP; 6 A; 4 C; 4 G; 10 T; 0 other;

Query Match 95.0%; Score 19; DB 24; Length 24;  
 Best Local Similarity 100.0%; Pred. No. 4.3;  
 Matches 19; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAGT 19  
 |||||  
 Db 6 TATGCATATTCCTGTAAGT 24

RESULT 4  
 AAT41604  
 ID AAT41604 standard; DNA; 19 BP.  
 XX  
 AC AAT41604;

DT 04-JUN-1997 (first entry)

DE Oligonucleotide containing core DNA regulatory element.

XX  
 KW Regulatory element; STAT; protein; cytokine; responsive;  
 KW host cell; transfection; agonist; antagonist; mediated; STAT5;  
 KW transcription; modulation; signalling pathway; STAT6;  
 KW oligonucleotide; electrophoretic mobility shift assay; EMSA; ds.

XX Synthetic.

XX  
 FH Key Location/Qualifiers  
 FT misc\_feature 5..19  
 FT /\*tag= a  
 FT /note= "core DNA regulatory element"

XX WO9630515-A1.

XX 03-OCT-1996.

XX 25-MAR-1996; 96WO-0504012.

XX 27-MAR-1995; 95US-0411020.

XX (LIGA-) LIGAND PHARM INC.

XX Lamb IP, Seidel HM, Tian Chan S;

XX WPI; 1996-455362/45.

XX DNA construct for screening modulators of cytokine-mediated  
 XX transcription - contg. regulatory element and a cytokine-sensitive  
 XX promoter operably linked to a heterologous gene

XX Example 1; Page 25; 72pp; English.

XX A novel DNA construct comprises the present oligonucleotide (ON),  
 CC which contains a core a regulatory element, operably linked to a  
 CC promoter, which is operably linked to a heterologous gene  
 CC (preferably a marker gene). The gene is under the transcriptional  
 CC control of the promoter and the ON sequence when the ON is bound by

CC a STAT protein activated in response to IL-2, IL-3, G-CSF, GM-CSF,  
 CC erythropoietin, thrombopoietin, or preferably IL-4, IL-7, IL-9,  
 CC IL-13 or IL-15. Cytokine responsive host cells transfected with the  
 CC DNA construct can be used to measure the ability of a compound to  
 CC act as an agonist or antagonist of cytokine mediated gene for  
 CC transcription. In particular, they can be used to screen for  
 CC cytokine modulators involved in the STAT5 and/or STAT6 protein  
 CC signalling pathway.  
 CC Following an electrophoretic mobility shift assay the DNA construct  
 CC was found to bind IL-4 weakly and IL-13 not determined.

XX Sequence 19 BP; 5 A; 3 C; 4 G; 7 T; 0 other;

Query Match 80.0%; Score 16; DB 17; Length 19;  
 Best Local Similarity 100.0%; Pred. No. 1.3e+02;  
 Matches 16; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 5 CATATTCCTGTAAGTG 20  
 |||||  
 Db 4 CATATTCCTGTAAGTG 19

RESULT 5  
 AAT31283  
 ID AAT31283 standard; DNA; 17 BP.

XX AAT31283;

XX 24-OCT-1996 (first entry)

XX STAT probe Ly6E.

XX  
 KW STAT; signal transducer and activator of transcription;  
 KW DNA binding protein; ligand; receptor; oncogenesis; inflammation;  
 KW autoimmune disease; antagonist; gene therapy; probe; ds.

XX Synthetic.

XX WO9620954-A2.

XX 11-JUL-1996.

XX 28-DEC-1995; 95WO-US17025.

XX 06-JAN-1995; 95US-0369796.

XX (UYRQ ) UNIV ROCKEFELLER.

XX Darnell JE, Horvath CM, Wen Z, Zhong Z;

XX WPI; 1996-333941/33.

XX New STAT protein DNA-binding domain peptide(s) - useful for  
 XX diagnosing, preventing or treating cellular dysfunction, e.g.  
 XX oncogenesis, inflammation, parasitic disease or autoimmunity

XX Example 1; Page 42; 138pp; English.

XX Synthetic DNA probes (AAT31281-86) were used in electrophoretic  
 XX mobility shift assays to determine the functionally active  
 XX regions of signal transducer and activator of transcription  
 XX (STAT) proteins STAT1 (see also AAW03176) and STAT3 (AAW03174).  
 XX Amino acids between approx. 400 and approx. 500 of these  
 XX human proteins determined DNA binding site specificity.  
 XX Mutations within this region resulted in greatly reduced DNA  
 XX binding affinities.

XX Sequence 17 BP; 5 A; 2 C; 3 G; 7 T; 0 other;

Query Match 75.0%; Score 15; DB 17; Length 17;  
 Best Local Similarity 100.0%; Pred. No. 4.2e+02;  
 Matches 15; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 6 ATATCTCTGTAAGT 20  
 Db 1 ATATCTCTGTAAGT 15

RESULT 6

AAQ69287/c  
 ID AAO69287 standard; DNA; 47 BP.  
 AC AAQ69287;  
 XX  
 XX 21-FEB-1995 (first entry)  
 DT  
 DE Human alpha-amylase gene, target region.  
 DE  
 XX DNA protein-binding assay; test sequence; screening sequence;  
 KW promoter; target; TATA box; Herpes Simplex Virus; HSV;  
 KW origin of replication; UL9; transcription factor; TFIID: ds.  
 XX  
 OS Synthetic.  
 XX  
 XX WO9414980-A.  
 PN  
 XX 07-JUL-1994.  
 PD  
 XX 20-DEC-1993; 93WO-US12388.  
 PF  
 XX 23-DEC-1992; 92US-0996783.  
 PR  
 XX 17-SEP-1993; 93US-0123936.  
 XX  
 PA (GENE-) GENELABS TECHNOLOGIES INC.  
 XX  
 XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI  
 XX WPI; 1994-234711/28.  
 DR  
 XX Sequence-directed DNA-binding molecules - useful in  
 PT pharmaceuticals and as molecular reagents  
 PT  
 XX Claim 28; Page 230; 587pp; English.

CC A DNA protein-binding assay is provided, useful for screening  
 CC libraries of synthetic or biological cpds. for their ability  
 CC to bind DNA test sequences. The assay is versatile in that any  
 CC number of test sequences can be tested by placing the test sequence  
 CC adjacent to a defined protein-binding screening sequence. Binding  
 CC of mols. to these test sequences changes the binding characteristics  
 CC of the protein mol. to its cognate binding sequence. When such a mol.  
 CC binds the test sequence, the equilibrium of the DNA:protein complexes  
 CC is disturbed, generating changes in the concentration of free DNA probe.  
 CC One application of this method is to eucaryotic general transcription  
 CC factors (e.g. TFIID), where the target region is typically selected  
 CC from DNA sequences adjacent to the binding site for the eucaryotic  
 CC transcription factor. Numerous exemplary test sequences are given:  
 CC the sequences in AAQ69251-731 and AAQ69850 correspond to promoter  
 CC targets (typically, TATA box-contg. sites) for human genes and the  
 CC sequences in AAQ69732-849 correspond to promoter targets for viral  
 CC genes.  
 CC The test sequences may also be randomly generated. DNA:protein  
 CC interaction may be used for screening purposes, e.g. the Herpes Simplex  
 CC virus (HSV) origin of replication and UL9 (see AAQ69851-52, AAQ69865 and  
 CC AAQ69891).

XX Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;

Query Match 74.0%; Score 14.8; DB 15; Length 47;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
 Db 28 TATTTATATTCCTGTAAG 11

RESULT 7

AAT63749/c  
 ID AAT63749 standard; DNA; 47 BP.  
 XX  
 AC AAT63749;  
 XX  
 XX 13-MAR-1997 (first entry)  
 DT  
 DE Human alpha amylase gene TFIID binding site.  
 DE  
 XX Duplex DNA; target region; binding characteristic; DNA binding protein;  
 KW TFIID; transcription factor; binding site; inhibition; enhance;  
 KW cancer; inherited genetic disorder; ds.  
 XX  
 OS Homo sapiens.  
 XX  
 XX US5578444-A.  
 PN  
 XX 26-NOV-1996.  
 PD  
 XX 27-JUN-1991; 91US-0723618.  
 PF  
 XX 20-DEC-1993; 93US-0171389.  
 PR  
 XX 27-JUN-1991; 91US-0723618.  
 PR  
 XX 23-DEC-1992; 92US-0996783.  
 PR  
 XX 17-SEP-1993; 93US-0123936.  
 XX  
 PA (GENE-) GENELABS TECHNOLOGIES INC.  
 XX  
 XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI  
 XX WPI; 1997-020402/02.  
 DR  
 XX Altering binding characteristics of DNA binding proteins to duplex  
 PT DNA - by attaching specific small cpd. to target region close to the  
 PT protein's binding site, useful in treatment of viral disease, cancer  
 PT etc  
 XX  
 XX Claim 6; Column 117; 264pp; English.

CC The sequences given in AAT63713-4312 represent duplex DNA's which act  
 CC as target regions in the method of the invention. The method for  
 CC altering the binding characteristics of a DNA-binding protein to duplex  
 CC DNA comprises contacting the duplex DNA with a small molecule which  
 CC binds sequence-specifically to a target region, where, when the small  
 CC molecule is bound to the target region, it is adjacent to, but not  
 CC overlapping by more than 4 bp, a binding site for a DNA-binding protein.  
 CC The small molecule is added at a concentration effective to alter the  
 CC binding of the DNA binding protein, pref. TFIID, to its binding site on  
 CC the duplex DNA. The binding of the small molecule may inhibit or  
 CC enhance the binding of the DNA-binding protein to its binding site. The  
 CC compounds isolated using this method are potentially useful as  
 CC therapeutic agents for treatment of any disease which involves a  
 CC specific DNA sequence, e.g. cancer, or inherited genetic disorders etc.  
 CC The method is suitable for screening large biological or chemical  
 CC libraries and allows determination of sequence-specific and relative  
 CC affinities of known DNA-binding agents for different DNA sequences.  
 CC The design of these duplex DNA's allows a single DNA:protein interaction  
 CC to be used for screening sequence-specific, or preferential, DNA binding  
 CC proteins that recognise almost any possible sequence (see also AAT49539-  
 CC 74).

XX Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;

Query Match 74.0%; Score 14.8; DB 18; Length 47;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18  
 Db 28 TATTTATATTCCTGTAAG 11

RESULT 8  
 AAX17037/c  
 ID AAX17037 standard; DNA; 47 BP.  
 AC  
 XX  
 AC AAX17037;  
 XX  
 DT 06-MAY-1999 (first entry)  
 XX  
 XX  
 DE Test sequence from human alpha-amylase gene.  
 XX  
 KW Test sequence; DNA-binding molecule; screening sequence; human;  
 KW nucleic acid amplification; target; viral; ds.  
 XX  
 KW Homo sapiens.  
 OS  
 XX US5869241-A.  
 PN  
 XX  
 PD 09-FEB-1999.  
 XX  
 XX 07-JUN-1995; 95US-0475228.  
 PF  
 XX 20-DEC-1993; 93US-0171389.  
 PR  
 XX 27-JUN-1991; 91US-0723618.  
 PR  
 XX 23-DEC-1992; 92US-0996783.  
 PR  
 XX 17-SEP-1993; 93US-0123936.  
 PR  
 XX 07-JUN-1995; 95US-0475228.  
 PR  
 XX  
 PA (GENE-) GENELABS TECHNOLOGIES INC.  
 XX  
 XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI  
 XX WPI; 1999-152755/13.  
 DR  
 XX  
 XX Determination of DNA sequence preference of a DNA-binding molecule -  
 PT based on inhibition of binding of protein to oligonucleotide  
 PT sequence attached to test sequence  
 PT  
 XX  
 PS Claim 3; Columns 119-120; 270pp; English.  
 XX  
 CC Sequences AAX17001 to AAX17600 represent specifically claimed target  
 CC test sequences that are used in the method of the invention of  
 CC determining the DNA sequence preference of a DNA-binding molecule. The  
 CC method comprises: (i) adding a test molecule and a DNA-binding protein to  
 CC a mixture of duplex DNA test oligonucleotides, each of the test  
 CC oligonucleotides having a test sequence adjacent to a screening sequence,  
 CC where the screening sequence binds to the DNA-binding protein with a  
 CC binding affinity that is independent of the DNA sequence of the test  
 CC sequence, and where the mixture of duplex DNA test oligonucleotides  
 CC includes several test sequences; (ii) incubating the test molecule, the  
 CC mixture of duplex DNA test oligonucleotides and the DNA-binding protein  
 CC for a time sufficient to permit binding of the test molecule to test  
 CC sequences in the duplex DNA; (iii) separating unbound test  
 CC oligonucleotides from test oligonucleotides bound to binding protein;  
 CC (iv) amplifying the unbound test oligonucleotides; (v) repeating steps  
 CC (ii) to (iv); (vi) isolating the amplified test oligonucleotides; and  
 CC (vii) sequencing the isolated test oligonucleotides. Test sequences  
 CC AAX17001-X17481 and AAX17600 correspond to promoter targets for human  
 CC genes and test sequences AAX17482-X17599 correspond to promoter targets  
 CC for viral genes.  
 XX  
 SQ Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;  
 Query Match 74.0%; Score 14.8; DB 20; Length 47;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 TATGCATATTCCTGTAAG 18  
 ||| |||||||||  
 Db 28 TATTATATTCCTGTAAG 11  
 RESULT 9  
 ABK82528/c  
 ID AAK69686 standard; DNA; 50 BP.  
 Query Match 74.0%; Score 14.8; DB 24; Length 47;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 TATGCATATTCCTGTAAG 18  
 ||| |||||||||  
 Db 28 TATTATATTCCTGTAAG 11  
 RESULT 10  
 AAQ69686/c  
 ID AAK69686 standard; DNA; 50 BP.  
 Query Match 74.0%; Score 14.8; DB 24; Length 47;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 TATGCATATTCCTGTAAG 18  
 ||| |||||||||  
 Db 28 TATTATATTCCTGTAAG 11

ID ABK82528 standard; DNA; 47 BP.  
 XX  
 AC ABK82528;  
 XX  
 DT 27-AUG-2002 (first entry)  
 XX  
 DE DNA binding molecule screening method test sequence #37.  
 XX  
 KW DNA binding molecule screening; inhibition of transcription;  
 KW infection; human immunodeficiency virus; HIV; parasite; cancer;  
 KW cardiovascular; respiratory; gastrointestinal; endocrine; metabolic;  
 KW rheumatic; immunological; haematological; neurological;  
 KW psychiatric; dermatological; ophthalmological; musculo-skeletal;  
 KW urogenital disorder; ss.  
 XX  
 OS Synthetic.  
 XX  
 PN US6384208-B1.  
 XX  
 PD 07-MAY-2002.  
 XX  
 XX 15-JUL-1999; 99US-0354947.  
 PF  
 XX 20-DEC-1993; 93US-0171389.  
 PR  
 XX 07-JUN-1995; 95US-0482080.  
 PR  
 XX 27-JUN-1991; 91US-0723618.  
 PR  
 XX 23-DEC-1992; 92US-0996783.  
 PR  
 XX 17-SEP-1993; 93US-0123936.  
 PR  
 XX  
 PA (GENE-) GENELABS TECHNOLOGIES INC.  
 XX  
 XX Edwards CA, Cantor CR, Andrews BM, Turin LM, Fry KE;  
 PI  
 XX WPI; 2002-442819/47.  
 DR  
 XX Decreasing transcriptional activity of genes for treating infections or  
 PT cancer, by administration of an agent that binds to two non-overlapping  
 PT regions of the gene -  
 PT  
 XX  
 PS Example 15; SEQ ID No 37; 98pp; English.  
 XX  
 CC The invention relates to a method of decreasing transcriptional activity  
 CC in a duplex deoxyribonucleic acid (DNA) template (T1) comprising  
 CC contacting (T1) with a binding agent comprising at least one small duplex  
 CC DNA-binding molecule (T2) coupled to at least one other small duplex-  
 CC binding molecule that binds to a non-overlapping region of target  
 CC sequence (T3). The method is useful for inhibiting transcription of a  
 CC range of disease-related genes for treating infections (by viruses,  
 CC including human immunodeficiency virus, bacteria, fungi, protozoa  
 CC and parasites), cancer, cardiovascular, respiratory, gastrointestinal,  
 CC endocrine/metabolic, rheumatic/immunological, haematological,  
 CC neurological, psychiatric, dermatological, ophthalmological,  
 CC musculo-skeletal, genetic or urogenital disorders. The method provides  
 CC sequence-specific inhibition of transcription of pathological genes  
 CC without affecting transcription of cellular genes regulated by the same  
 CC transcription factor, and can be applied to regulation of any gene.  
 CC ABK82492-ABK83155 represent DNA binding molecule test sequences used in  
 CC the method of the invention.  
 XX  
 SQ Sequence 47 BP; 17 A; 6 C; 10 G; 14 T; 0 other;  
 Query Match 74.0%; Score 14.8; DB 24; Length 47;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 TATGCATATTCCTGTAAG 18  
 ||| |||||||||  
 Db 28 TATTATATTCCTGTAAG 11  
 RESULT 10  
 AAQ69686/c  
 ID AAK69686 standard; DNA; 50 BP.  
 Query Match 74.0%; Score 14.8; DB 24; Length 47;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 TATGCATATTCCTGTAAG 18  
 ||| |||||||||  
 Db 28 TATTATATTCCTGTAAG 11

XX AAQ69686;  
 AC 02-MAR-1995 (first entry)  
 DT  
 XX Human pancreatic alpha-amylase gene, target region.  
 DE  
 XX  
 XX DNA protein-binding assay; test sequence; screening sequence;  
 KW promoter; target; TATA box; Herpes Simplex Virus; HSV;  
 KW origin of replication; UL9; transcription factor; TFIID: ds.  
 XX  
 XX Synthetic.  
 OS  
 XX WO9414980-A.  
 PN  
 XX 07-JUL-1994.  
 PD  
 XX 20-DEC-1993; 93WO-US12388.  
 PF  
 XX 23-DEC-1992; 92US-0996783.  
 PR  
 XX 17-SEP-1993; 93US-0123936.  
 PP  
 XX (GENE-) GENELABS TECHNOLOGIES INC.  
 PA  
 XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI WPI; 1994-234711/28.  
 XX  
 XX Sequence-directed DNA-binding molecules - useful in  
 PT pharmaceuticals and as molecular reagents  
 PS Claim 28; Page 430; 587pp; English.  
 XX  
 CC A DNA protein-binding assay is provided, useful for screening  
 CC libraries of synthetic or biological cpds. for their ability  
 CC to bind DNA test sequences. The assay is versatile in that any  
 CC number of test sequences can be tested by placing the test sequence  
 CC adjacent to a defined protein-binding screening sequence. Binding  
 CC of mols. to these test sequences changes the binding characteristics  
 CC of the protein mol. to its cognate binding sequence. When such a mol.  
 CC binds the test sequence, the equilibrium of the DNA:protein complexes  
 CC is disturbed, generating changes in the concentration of free DNA probe.  
 CC One application of this method is to eucaryotic general transcription  
 CC factors (e.g. TFIID), where the target region is typically selected  
 CC from DNA sequences adjacent to the binding site for the eucaryotic  
 CC transcription factor. Numerous exemplary test sequences are given:  
 CC the sequences in AAQ69251-731 and AAQ69850 correspond to promoter  
 CC targets (typically, TATA box-contg. sites) for human genes and the  
 CC sequences in AAQ69732-849 correspond to promoter targets for viral  
 CC genes. The test sequences may also be randomly generated. DNA:protein  
 CC interaction may be used for screening purposes, e.g. the Herpes Simplex  
 CC Virus (HSV) origin of replication and UL9 (see AAQ69851-52, AAQ69865 and  
 CC AAQ69891).  
 XX  
 SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;  
 Query Match 74.0%; Score 14.8; DB 15; Length 50;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 TATGCATATTCCTGTAAG 18  
 III |  
 DB 32 TATTATATTCCTGTAAG 15  
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 AAQ69687/c  
 ID AAQ69687 standard; DNA: 50 BP.  
 AC  
 XX AAQ69687;  
 DT 02-MAR-1995 (first entry)  
 XX

DE  
 XX Human pancreatic alpha-amylase gene, target region.  
 KW DNA protein-binding assay; test sequence; screening sequence;  
 KW promoter; target; TATA box; Herpes Simplex Virus; HSV;  
 KW origin of replication; UL9; transcription factor; TFIID: ds.  
 XX  
 XX Synthetic.  
 OS  
 XX WO9414980-A.  
 PN  
 XX 07-JUL-1994.  
 PD  
 XX 20-DEC-1993; 93WO-US12388.  
 PF  
 XX 23-DEC-1992; 92US-0996783.  
 PR  
 XX 17-SEP-1993; 93US-0123936.  
 PP  
 XX (GENE-) GENELABS TECHNOLOGIES INC.  
 PA  
 XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;  
 PI WPI; 1994-234711/28.  
 XX  
 XX Sequence-directed DNA-binding molecules - useful in  
 PT pharmaceuticals and as molecular reagents  
 PS Claim 28; Page 430; 587pp; English.  
 XX  
 CC A DNA protein-binding assay is provided, useful for screening  
 CC libraries of synthetic or biological cpds. for their ability  
 CC to bind DNA test sequences. The assay is versatile in that any  
 CC number of test sequences can be tested by placing the test sequence  
 CC adjacent to a defined protein-binding screening sequence. Binding  
 CC of mols. to these test sequences changes the binding characteristics  
 CC of the protein mol. to its cognate binding sequence. When such a mol.  
 CC binds the test sequence, the equilibrium of the DNA:protein complexes  
 CC is disturbed, generating changes in the concentration of free DNA probe.  
 CC One application of this method is to eucaryotic general transcription  
 CC factors (e.g. TFIID), where the target region is typically selected  
 CC from DNA sequences adjacent to the binding site for the eucaryotic  
 CC transcription factor. Numerous exemplary test sequences are given:  
 CC the sequences in AAQ69251-731 and AAQ69850 correspond to promoter  
 CC targets (typically, TATA box-contg. sites) for human genes and the  
 CC sequences in AAQ69732-849 correspond to promoter targets for viral  
 CC genes. The test sequences may also be randomly generated. DNA:protein  
 CC interaction may be used for screening purposes, e.g. the Herpes Simplex  
 CC Virus (HSV) origin of replication and UL9 (see AAQ69851-52, AAQ69865 and  
 CC AAQ69891).  
 XX  
 SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;  
 Query Match 74.0%; Score 14.8; DB 15; Length 50;  
 Best Local Similarity 88.9%; Pred. No. 5.9e+02;  
 Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;  
 QY 1 TATGCATATTCCTGTAAG 18  
 III |  
 DB 32 TATTATATTCCTGTAAG 15  
 RESULT 12  
 AAQ64148/c  
 ID AAT64148 standard; DNA: 50 BP.  
 XX  
 AC AAT64148;  
 XX  
 XX 17-MAR-1997 (first entry)  
 DT  
 XX Human pancreatic alpha-amylase gene TFIID binding site.  
 DE  
 XX Duplex DNA: target region; binding characteristic; DNA binding protein;  
 KW TFIID; transcription factor; binding site; inhibition; enhance;  
 KW cancer; inherited genetic disorder; ds.

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XX OS Homo sapiens.
XX PN US5578444-A.
XX PD 26-NOV-1996.
XX PF 27-JUN-1991; 91US-0723618.
XX PR 20-DEC-1993; 93US-0171389.
XX PR 27-JUN-1991; 91US-0723618.
XX PR 23-DEC-1992; 92US-0996783.
XX PR 17-SEP-1993; 93US-0123936.
XX PA (GENE-) GENELABS TECHNOLOGIES INC.
XX PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX WPI: 1997-020402/02.
XX
PT Altering binding characteristics of DNA binding proteins to duplex
PT DNA - by attaching specific small cpd. to target region close to the
PT protein's binding site, useful in treatment of viral disease, cancer
PT etc
XX
XX Claim 6; Column 321-322; 264pp; English.
XX
CC The sequences given in AAT63713-4312 represent duplex DNA's which act
CC as target regions in the method of the invention. The method for
CC altering the binding characteristics of a DNA-binding protein to duplex
CC DNA comprises contacting the duplex DNA with a small molecule which
CC binds sequence-specifically to a target region, where, when the small
CC molecule is bound to the target region, it is adjacent to, but not
CC overlapping by more than 4 bp, a binding site for a DNA-binding protein.
CC The small molecule is added at a concentration effective to alter the
CC binding of the DNA binding protein, pref. TFIID, to its binding site on
CC the duplex DNA. The binding of the small molecule may inhibit or
CC enhance the binding of the DNA-binding protein to its binding site. The
CC compounds isolated using this method are potentially useful as
CC therapeutic agents for treatment of any disease which involves a
CC specific DNA sequence, e.g. cancer, or inherited genetic disorders etc.
CC The method is suitable for screening large biological or chemical
CC libraries and allows determination of sequence-specific and relative
CC affinities of known DNA-binding agents for different DNA sequences.
CC The design of these duplex DNA's allows a single DNA:protein interaction
CC to be used for screening sequence-specific, or preferential, DNA binding
CC proteins that recognise almost any possible sequence (see also AAT49539-
CC 74).
XX
SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
Query Match 74.0%; Score 14.8; DB 18; Length 50;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
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DB 32 TATTTATATTCCTGTAAG 15

RESULT 13
AAT64149/c
ID AAT64149 standard; DNA; 50 BP.
XX
AC AAT64149;
XX
DT 17-MAR-1997 (first entry)
XX
DE Human pancreatic amylase gene TFIID binding site.
XX
KW Duplex DNA; target region; binding characteristic; DNA binding protein;
KW TFIID; transcription factor; binding site; inhibition; enhance;
KW cancer; inherited genetic disorder; ds.

XX OS Homo sapiens.
XX PN US5578444-A.
XX PD 26-NOV-1996.
XX PF 27-JUN-1991; 91US-0723618.
XX PR 20-DEC-1993; 93US-0171389.
XX PR 27-JUN-1991; 91US-0723618.
XX PR 23-DEC-1992; 92US-0996783.
XX PR 17-SEP-1993; 93US-0123936.
XX PA (GENE-) GENELABS TECHNOLOGIES INC.
XX PI Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX WPI: 1997-020402/02.
XX
PT Altering binding characteristics of DNA binding proteins to duplex
PT DNA - by attaching specific small cpd. to target region close to the
PT protein's binding site, useful in treatment of viral disease, cancer
PT etc
XX
XX Claim 6; Column 321-322; 264pp; English.
XX
CC The sequences given in AAT63713-4312 represent duplex DNA's which act
CC as target regions in the method of the invention. The method for
CC altering the binding characteristics of a DNA-binding protein to duplex
CC DNA comprises contacting the duplex DNA with a small molecule which
CC binds sequence-specifically to a target region, where, when the small
CC molecule is bound to the target region, it is adjacent to, but not
CC overlapping by more than 4 bp, a binding site for a DNA-binding protein.
CC The small molecule is added at a concentration effective to alter the
CC binding of the DNA binding protein, pref. TFIID, to its binding site on
CC the duplex DNA. The binding of the small molecule may inhibit or
CC enhance the binding of the DNA-binding protein to its binding site. The
CC compounds isolated using this method are potentially useful as
CC therapeutic agents for treatment of any disease which involves a
CC specific DNA sequence, e.g. cancer, or inherited genetic disorders etc.
CC The method is suitable for screening large biological or chemical
CC libraries and allows determination of sequence-specific and relative
CC affinities of known DNA-binding agents for different DNA sequences.
CC The design of these duplex DNA's allows a single DNA:protein interaction
CC to be used for screening sequence-specific, or preferential, DNA binding
CC proteins that recognise almost any possible sequence (see also AAT49539-
CC 74).
XX
SQ Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
Query Match 74.0%; Score 14.8; DB 18; Length 50;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 TATGCATATTCCTGTAAG 18
   ||| |||||
DB 32 TATTTATATTCCTGTAAG 15

RESULT 14
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ID AAX17436 standard; DNA; 50 BP.
XX
AC AAX17436;
XX
DT 06-MAY-1999 (first entry)
XX
DE Test sequence from human pancreatic alpha-amylase gene.
XX
KW Test sequence; DNA-binding molecule; screening sequence; human;
KW nucleic acid amplification; target; viral; ds.

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OS Homo sapiens.
XX US5869241-A.
XX 09-FEB-1999.
XX 07-JUN-1995; 950S-0475228.
XX 20-DEC-1993; 930S-0171389.
XX 27-JUN-1991; 910S-0723618.
XX 23-DEC-1992; 920S-0996783.
XX 17-SEP-1993; 930S-0123936.
XX 07-JUN-1995; 950S-0475228.
XX (GENE-) GENELABS TECHNOLOGIES INC.
XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX WPI; 1999-152755/13.
XX Determination of DNA sequence preference of a DNA-binding molecule -
XX based on inhibition of binding of protein to oligonucleotide
XX sequence attached to test sequence
XX Claim 3; Columns 323-324; 270pp: English.
XX Sequences AAX17001 to AAX17600 represent specifically claimed target
XX test sequences that are used in the method of the invention of
XX determining the DNA sequence preference of a DNA-binding molecule. The
XX method comprises: (i) adding a test molecule and a DNA-binding protein to
XX a mixture of duplex DNA test oligonucleotides, each of the test
XX oligonucleotides having a test sequence adjacent to a screening sequence,
XX where the screening sequence binds to the DNA-binding protein with a
XX binding affinity that is independent of the DNA sequence of the test
XX sequence, and where the mixture of duplex DNA test oligonucleotides
XX includes several test sequences; (ii) incubating the test molecule, the
XX mixture of duplex DNA test oligonucleotides and the DNA-binding protein
XX for a time sufficient to permit binding of the test molecule to test
XX sequences in the duplex DNA; (iii) separating unbound test
XX oligonucleotides from test oligonucleotides bound to binding protein;
XX (iv) amplifying the unbound test oligonucleotides; (v) repeating steps
XX (ii) to (iv); (vi) isolating the amplified test oligonucleotides; and
XX (vii) sequencing the isolated test oligonucleotides. Test sequences
XX AAX17001-X17481 and AAX17600 correspond to promoter targets for human
XX genes and test sequences AAX17482-X17599 correspond to promoter targets
XX for viral genes.
XX Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
SQ
Query Match 74.0%; Score 14.8; DB 20; Length 50;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
Matches 16; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 TATGCATATTCCTGTAAG 18
DB 32 TATTTATATTCCTGTAAG 15
RESULT 15
AAX17437/C
XX AAX17437 standard; DNA; 50 BP.
XX AC AAX17437;
XX DT 06-MAY-1999 (first entry)
XX DE Test sequence from human pancreatic amylase gene.
XX KW Test sequence; DNA-binding molecule; screening sequence; human;
XX KW nucleic acid amplification; target; viral; ds.
XX OS Homo sapiens.

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PN US5869241-A.
XX 09-FEB-1999.
XX 07-JUN-1995; 950S-0475228.
XX 20-DEC-1993; 930S-0171389.
XX 27-JUN-1991; 910S-0723618.
XX 23-DEC-1992; 920S-0996783.
XX 17-SEP-1993; 930S-0123936.
XX 07-JUN-1995; 950S-0475228.
XX (GENE-) GENELABS TECHNOLOGIES INC.
XX Andrews BM, Cantor CR, Edwards CA, Fry KE, Turin LM;
XX WPI; 1999-152755/13.
XX Determination of DNA sequence preference of a DNA-binding molecule -
XX based on inhibition of binding of protein to oligonucleotide
XX sequence attached to test sequence
XX Claim 3; Columns 323-324; 270pp: English.
XX Sequences AAX17001 to AAX17600 represent specifically claimed target
XX test sequences that are used in the method of the invention of
XX determining the DNA sequence preference of a DNA-binding molecule. The
XX method comprises: (i) adding a test molecule and a DNA-binding protein to
XX a mixture of duplex DNA test oligonucleotides, each of the test
XX oligonucleotides having a test sequence adjacent to a screening sequence,
XX where the screening sequence binds to the DNA-binding protein with a
XX binding affinity that is independent of the DNA sequence of the test
XX sequence, and where the mixture of duplex DNA test oligonucleotides
XX includes several test sequences; (ii) incubating the test molecule, the
XX mixture of duplex DNA test oligonucleotides and the DNA-binding protein
XX for a time sufficient to permit binding of the test molecule to test
XX sequences in the duplex DNA; (iii) separating unbound test
XX oligonucleotides from test oligonucleotides bound to binding protein;
XX (iv) amplifying the unbound test oligonucleotides; (v) repeating steps
XX (ii) to (iv); (vi) isolating the amplified test oligonucleotides; and
XX (vii) sequencing the isolated test oligonucleotides. Test sequences
XX AAX17001-X17481 and AAX17600 correspond to promoter targets for human
XX genes and test sequences AAX17482-X17599 correspond to promoter targets
XX for viral genes.
XX Sequence 50 BP; 18 A; 6 C; 10 G; 16 T; 0 other;
SQ
Query Match 74.0%; Score 14.8; DB 20; Length 50;
Best Local Similarity 88.9%; Pred. No. 5.9e+02;
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DB 32 TATTTATATTCCTGTAAG 15
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GenCore version 5.1.3  
Copyright (c) 1993 - 2002 CompuGen Ltd.

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Searched: 2054640 segs, 14551402878 residues

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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

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- 13: gb\_un.\*
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- 41: em\_htgo\_other.\*

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

SUMMARIES

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2	20	100.0	20	6	A90874	A90874 Sequence 9
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4	20	100.0	20	6	AX455575	AX455575 Sequence
5	20	100.0	27	6	AR063217	AR063217 Sequence
6	20	100.0	27	6	AX001643	AX001643 Sequence
7	20	100.0	27	6	I49528	I49528 Sequence 2
8	19	95.0	20	6	A89784	A89784 Sequence 6
9	19	95.0	20	6	A90871	A90871 Sequence 6
10	16.4	82.0	32	6	AR135319	AR135319 Sequence
11	16.4	82.0	50	6	AR032839	AR032839 Sequence
12	16.4	82.0	50	6	AR209503	AR209503 Sequence
13	16.4	82.0	50	6	I29579	I29579 Sequence 45
14	16.4	82.0	50	6	I91253	I91253 Sequence 45
15	15.8	79.0	20	6	A89786	A89786 Sequence 8
16	15.8	79.0	20	6	A90873	A90873 Sequence 8
17	14.4	72.0	83	6	AX328437	AX328437 Sequence
18	14.2	71.0	41	6	AR109119	AR109119 Sequence
19	14.2	71.0	41	6	AR200774	AR200774 Sequence
20	13.8	69.0	71	6	AR122803	AR122803 Sequence
21	13.6	68.0	51	6	AX162422	AX162422 Sequence
22	13.2	66.0	25	6	AX117480	AX117480 Sequence
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24	13.2	66.0	64	9	AF032251	AF032251 Otolomur
25	13.2	66.0	75	9	D86841S01	D86841 Homo sapien
26	13.2	66.0	76	9	AF032248	AF032248 Otolomur
27	13.2	66.0	76	9	AF032254	AF032254 Otolomur
28	13	65.0	26	6	A89779	A89779 Sequence 1
29	13	65.0	26	6	A90893	A90893 Sequence 28
30	12.8	64.0	21	6	A91594	A91594 Sequence 12
31	12.8	64.0	51	6	AX159923	AX159923 Sequence
32	12.8	64.0	51	6	AX159925	AX159925 Sequence
33	12.8	64.0	51	6	AX159926	AX159926 Sequence
34	12.8	64.0	51	6	AX160807	AX160807 Sequence
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ALIGNMENTS

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SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL

A89787  
Sequence 9 from Patent WO9832462.  
A89787  
A89787.1 GI:6738301  
unidentified.  
unclassified.  
1 (bases 1 to 20)  
Lipford,G.B. and Heeg,K.  
PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
Patent: WO 9832462-A 9 30-JUL-1998;

20 bp  
DNA  
linear  
PAT 22-JAN-2000

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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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VERSION     A90874.1 GI:6739268
KEYWORDS   unidentified.
SOURCE      unidentified.
ORGANISM    unidentified.
REFERENCE   1 (bases 1 to 20)
AUTHORS     Heeg,K.P. and Lipford,G.B.
TITLE       Pharmaceutical composition comprising a polynucleotide and an
            antigen especially for vaccination
JOURNAL     Patent: EP 0855184-A 9 29-JUL-1998;
            HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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ACCESSION  AR078393
VERSION     AR078393.1 GI:10005139
KEYWORDS   Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 20)
AUTHORS     Bachmaier,K., Hessel,A.,John., Neu,N. and Penninger,J.Martin.
TITLE       Peptides capable of modulating inflammatory heart disease
JOURNAL     Patent: US 5962636-A 10 05-OCT-1999;
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ACCESSION  AX455575
VERSION     AX455575.1 GI:21714643
KEYWORDS   synthetic construct.
SOURCE      synthetic construct.
ORGANISM    artificial sequences.
REFERENCE   1
AUTHORS     Bauer,S., Lipford,G. and Wagner,H.
TITLE       Process for high throughput screening of cpq-based
            immuno-agonist/antagonist
JOURNAL     Patent: WO 0222809-A 52 21-MAR-2002;
            Coley Pharmaceutical GmbH (DE)
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VERSION     AR063217.1 GI:5990908
KEYWORDS   Unknown.
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE   1 (bases 1 to 27)
AUTHORS     Hinrichs,S.H. and Orten,D.Jo.
TITLE       Methods for inhibiting transcription of the cyclic AMP responsive
            element binding protein and the activating transcription factor 1
JOURNAL     Patent: US 5844096-A 2 01-DEC-1998;
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ACCESSION  AX001643

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VERSION AX001643.1 GI:7241772  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unidentified  
unclassified.  
REFERENCE 1 (bases 1 to 27)  
AUTHORS Damert,A. and Plate,K.  
TITLE REGULATORY SEQUENCES INVOLVED IN HYPOXIA REGULATED GENE EXPRESSION  
AND USES THEREOF  
JOURNAL Patent: WO 9856936-A 15 17-DEC-1998;  
MAX PLANCK GESFELLSCHAFT (DE); DAMERT ANNETTE (DE)  
FEATURES  
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ACCESSION 149528  
VERSION 149528.1 GI:2471748  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 27)  
AUTHORS Hintichs,S.H. and Orten,D.Jo.  
TITLE Methods for inhibiting transcription of the cyclic AMP responsive  
element binding protein and the activating transcription factor 1  
JOURNAL Patent: US 5641486-A 2 24-JUN-1997;  
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ACCESSION A89784  
VERSION A89784.1 GI:6738298  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Lipford,G.B. and Heeg,K.  
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
JOURNAL Patent: WO 9832462-A 6 30-JUL-1998;  
LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)

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ACCESSION A90871  
VERSION A90871.1 GI:6739265  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Heeg,K.P. and Lipford,G.B.  
TITLE Pharmaceutical composition comprising a polynucleotide and an  
antigen especially for vaccination  
JOURNAL Patent: EP 0855184-A 6 29-JUL-1998;  
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)  
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ACCESSION ARL35319  
VERSION ARL35319.1 GI:14124224  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 32)  
AUTHORS Leiden,J.M.  
TITLE Mouse model for congestive heart failure  
JOURNAL Patent: US 6194632-A 1 27-FEB-2001;  
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GenCore version 5.1.3  
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OM nucleic - nucleic search, using sw model

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Perfect score: 20

Sequence: 1 gtatttccagaaaaaggaac 20

Scoring table: IDENTITY\_NUC

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Total number of hits satisfying chosen parameters: 995600.

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Listing first 45 summaries

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4	20	100.0	20	6	AR204918	AR204918 Sequence
5	20	100.0	20	6	AR204923	AR204923 Sequence
6	20	100.0	20	6	AX455586	AX455586 Sequence
7	20	100.0	20	6	I33466	I33466 Sequence 3
8	20	100.0	20	6	I40053	I40053 Sequence 3
9	20	100.0	20	6	I47062	I47062 Sequence 3
10	20	100.0	20	6	I81487	I81487 Sequence 3
11	20	100.0	30	6	AR182577	AR182577 Sequence 3
12	20	100.0	42	6	A37863	A37863 Sequence 6
13	20	100.0	100	6	AR174601	AR174601 Sequence
14	20	100.0	100	6	AR174602	AR174602 Sequence
15	20	100.0	100	6	AX047032	AX047032 Sequence
16	20	100.0	100	6	AX047033	AX047033 Sequence
17	20	100.0	100	6	AX280202	AX280202 Sequence
18	20	100.0	100	6	AX280203	AX280203 Sequence
19	20	100.0	100	6	AX365191	AX365191 Sequence
20	20	100.0	100	6	AX365192	AX365192 Sequence
21	17	85.0	17	6	AR165233	AR165233 Sequence
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37	14.2	71.0	69	3	AF250305	AF250305 Chironomu
38	14.2	71.0	98	6	AX354846	AX354846 Sequence
39	14	70.0	16	6	AR165232	AR165232 Sequence
40	14	70.0	16	6	AR201403	AR201403 Sequence
41	14	70.0	16	6	AX338643	AX338643 Sequence
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43	14	70.0	16	6	I15311	I15311 Sequence 12
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DEFINITION  
ACCESSION  
VERSION  
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SOURCE  
ORGANISM  
REFERENCE  
AUTHORS  
TITLE  
JOURNAL

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Sequence 22 from Patent WO9832462.  
A89800  
A89800.1 GI:6738314  
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unclassified.  
1 (bases 1 to 20)  
Lipford,G.B. and Heeg,K.  
PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
Patent: WO 9832462-A 22 30-JUL-1998;  
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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)
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REFERENCE  1 (bases 1 to 20)
AUTHORS   Heeg, K.P. and Lipford, G.B.
TITLE     Pharmaceutical composition comprising a polynucleotide and an
          antigen especially for vaccination
JOURNAL   Patent: EP 0855184-A 22 29-JUL-1998;
          HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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VERSION     AR010171.1 GI:3968976
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SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 20)
AUTHORS   Hoey, T. and Rothe, M.
TITLE     Nucleic acid encoding human signal transducer and activator of
          transcription 4
JOURNAL   Patent: US 5756700-A 3 26-MAY-1998;
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ACCESSION  AR204918
VERSION     AR204918.1 GI:21502366
KEYWORDS   Unknown.
SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 20)
AUTHORS   LaRoche, W., Patel, B.K.R. and Pierce, J.H.
TITLE     Attenuated and dominant negative variant cDNAs of Stat6b and
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JOURNAL   Patent: US 6368828-A 9 09-APR-2002;
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VERSION     AR204923.1 GI:21502372
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SOURCE     Unknown.
ORGANISM   Unclassified.
REFERENCE  1 (bases 1 to 20)
AUTHORS   LaRoche, W., Patel, B.K.R. and Pierce, J.H.
TITLE     Attenuated and dominant negative variant cDNAs of Stat6: Stat6b and
          Stat6c
JOURNAL   Patent: US 6368828-A 14 09-APR-2002;
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VERSION     AX455586.1 GI:21714654
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AUTHORS   LaRoche, W., Patel, B.K.R. and Pierce, J.H.
TITLE     Attenuated and dominant negative variant cDNAs of Stat6: Stat6b and
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JOURNAL   Patent: US 6368828-A 14 09-APR-2002;
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TITLE     Attenuated and dominant negative variant cDNAs of Stat6: Stat6b and
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ORGANISM    synthetic construct
artificial sequences.
REFERENCE   1
AUTHORS     Bauer,S., Lipford,G. and Wagner,H.
TITLE       Process for high throughput screening of cpv-based
            immuno-agonist/antagonist
JOURNAL     Patent: WO 0222809-A 63 21-MAR-2002;
            Coley Pharmaceutical GmbH (DE)
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VERSION   I33466.1 GI:1824257
KEYWORDS  .
SOURCE    Unknown.
ORGANISM  Unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS   McKnight,S.L. and Hou,J.
TITLE     Interleukin 4 signal transducers
JOURNAL   Patent: US 5591825-A 3 07-JAN-1997;
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KEYWORDS  .
SOURCE    Unknown.
ORGANISM  Unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS   McKnight,S.L., Hou,J. and Schindler,U.
TITLE     Interleukin-2 signal transducers and binding assays
JOURNAL   Patent: US 5618693-A 3 08-APR-1997;
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artificial sequences.
REFERENCE   1
AUTHORS     Bauer,S., Lipford,G. and Wagner,H.
TITLE       Process for high throughput screening of cpv-based
            immuno-agonist/antagonist
JOURNAL     Patent: WO 0222809-A 63 21-MAR-2002;
            Coley Pharmaceutical GmbH (DE)
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KEYWORDS  .
SOURCE    Unknown.
ORGANISM  Unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS   Hoeft,T. and Rothe,M.
TITLE     Human signal transducer and binding assays
JOURNAL   Patent: US 5639858-A 3 17-JUN-1997;
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VERSION   I81487.1 GI:3209784
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SOURCE    Unknown.
ORGANISM  Unclassified.
REFERENCE 1 (bases 1 to 20)
AUTHORS   McKnight,S.L. and Hou,J.
TITLE     Nucleic acid encoding an interleukin 4 signal transducer
JOURNAL   Patent: US 5710266-A 3 20-JAN-1998;
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RESULT 11
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DEFINITION Sequence 25 from patent US 6338949.

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ACCESSION AR182577  
VERSION AR182577.1 GI:20225784  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 30)  
AUTHORS Darnell, J.E. Jr., Schindler, C.W., Fu, X.-Y., Wen, Z. and Zhong, Z.  
TITLE Nucleic acids encoding receptor recognition factor stat4 and methods of use thereof  
JOURNAL Patent: US 638949-A 25 15-JAN-2002;  
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ACCESSION A37863  
VERSION A37863.1 GI:2294543  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 42)  
AUTHORS Benach, P., Perez, C. and Wietzerbin, J.  
TITLE DNA SEQUENCES INVOLVED IN THE TRANSCRIPTION OF GENES UNDER THE EFFECT OF INDUCERS, AND BIOLOGICAL APPLICATIONS THEREOF  
JOURNAL Patent: WO 9408025-A 6 14-APR-1994;  
COMMENT INST NAT SANTE RECH MED (FR)  
OTHER PUBLICATION FR 2696181 940401.  
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ACCESSION AR174601  
VERSION AR174601.1 GI:17914921  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 100)  
AUTHORS Novak, J.E., Presnell, S.R., Sprecher, C.A., Foster, D.C., Holly, R.D., Gross, J.A., Johnston, J.V., Nelson, A.J., Dillon, S.R. and Hammond, A.K.

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Db 6 GTATTCCCGAAGGAAC 25

RESULT 14  
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ACCESSION AR174602  
VERSION AR174602.1 GI:17914922  
KEYWORDS  
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ORGANISM Unknown.  
REFERENCE 1 (bases 1 to 100)  
AUTHORS Novak, J.E., Presnell, S.R., Sprecher, C.A., Foster, D.C., Holly, R.D., Gross, J.A., Johnston, J.V., Nelson, A.J., Dillon, S.R. and Hammond, A.K.  
TITLE Cytokine zalphall Ligand  
JOURNAL Patent: US 6307024-A 60 23-OCT-2001;  
FEATURES Location/Qualifiers  
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Db 44 GTATTCCCGAAGGAAC 63

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AX047032  
LOCUS AX047032  
DEFINITION Sequence 37 from Patent WO0068381.  
ACCESSION AX047032  
VERSION AX047032.1 GI:11876456  
KEYWORDS  
SOURCE synthetic construct.  
ORGANISM synthetic construct.  
REFERENCE 1 (bases 1 to 100)  
AUTHORS Presnell, S.R., Foster, D.C., Hammond, A.K. and Lok, S.  
TITLE Cytokine receptor mouse zcytor10  
JOURNAL Patent: WO 0068381-A 37 16-NOV-2000;  
ZymoGenetics, Inc. (US)  
FEATURES Location/Qualifiers  
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Search completed: December 12, 2002, 02:56:14  
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Copyright (c) 1993 - 2002 Compugen Ltd.

OM nucleic - nucleic search, using sw model  
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(without alignments)  
1829.698 Million cell updates/sec

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Sequence: 1 agatttctagaattcaatc 20

Scoring table: IDENTITY\_NUC  
Gapop 10.0 , Gapext 1.0

Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0  
Maximum DB seq length: 100

Post-processing: Minimum Match 0%  
Maximum Match 100%  
Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a

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and is derived by analysis of the total score distribution.

## SUMMARIES

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3	20	100.0	20	6	AX455553	AX455553 Sequence
4	20	100.0	21	6	I23456	I23456 Sequence 8
5	20	100.0	100	6	ARI74601	ARI74601 Sequence
c 6	20	100.0	100	6	ARI74602	ARI74602 Sequence
c 7	20	100.0	100	6	AX047032	AX047032 Sequence
c 8	20	100.0	100	6	AX047033	AX047033 Sequence
c 9	20	100.0	100	6	AX280202	AX280202 Sequence
c 10	20	100.0	100	6	AX280203	AX280203 Sequence
c 11	20	100.0	100	6	AX365191	AX365191 Sequence
c 12	20	100.0	100	6	AX365192	AX365192 Sequence
c 13	14.2	71.0	31	6	ARI19638	ARI19638 Sequence
c 14	14.2	71.0	31	6	ARI169119	ARI169119 Sequence
c 15	14.2	71.0	58	6	ARI138229	ARI138229 Sequence
c 16	14.2	71.0	64	12	SYNCOCK	K01192 Yeast (S.ce
c 17	13.8	69.0	24	6	A03717	A03717 Oligonucleo
c 18	13.8	69.0	41	6	AX343814	AX343814 Sequence
c 19	13.8	69.0	41	6	AX343816	AX343816 Sequence
c 20	13.8	69.0	44	6	AX343812	AX343812 Sequence
c 21	13.8	69.0	44	6	AX343818	AX343818 Sequence
c 22	13.8	69.0	76	8	YSCGNC152	M87429 Yeast Eco R
c 23	13.8	69.0	91	1	AF087321	AF087321 Chlamydia
c 24	13.8	69.0	94	10	AY041972	AY041972 Phodopus
c 25	13.6	68.0	25	6	AR043941	AR043941 Sequence
c 26	13.6	68.0	25	6	AR073474	AR073474 Sequence
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c 28	13.6	68.0	25	6	I93345	I93345 Sequence 15
c 29	13.6	68.0	26	6	AR043992	AR043992 Sequence
c 30	13.6	68.0	26	6	AR073525	AR073525 Sequence
c 31	13.6	68.0	26	6	AX022135	AX022135 Sequence
c 32	13.6	68.0	26	6	I93396	I93396 Sequence 82
c 33	13.6	68.0	65	6	AX483015	AX483015 Sequence
c 34	13.6	68.0	72	6	AR043980	AR043980 Sequence
c 35	13.6	68.0	72	6	AR073513	AR073513 Sequence
c 36	13.6	68.0	72	6	AX022123	AX022123 Sequence
c 37	13.6	68.0	72	6	I93384	I93384 Sequence 70
c 38	13.2	66.0	31	6	AX327657	AX327657 Sequence
c 39	13.2	66.0	45	6	AR076832	AR076832 Sequence
c 40	13.2	66.0	51	6	AX160107	AX160107 Sequence
c 41	13.2	66.0	58	6	ARI40194	ARI40194 Sequence
c 42	13.2	66.0	58	6	ARI73275	ARI73275 Sequence
c 43	13.2	66.0	63	6	ARI138221	ARI138221 Sequence
c 44	13.2	66.0	72	6	I74780	I74780 Sequence 12
c 45	13.2	66.0	74	10	RATPAM25	U52661 Rattus norv

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## RESULT 1

A89799 A89799 Sequence 21 from Patent WO9832462. 20 bp DNA linear PAT 22-JAN-2000  
LOCUS  
DEFINITION  
ACCESSION A89799  
VERSION A89799.1 GI:6738313  
KEYWORDS  
SOURCE unidentified.  
ORGANISM unclassified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Lipford G.B. and Heeg K.  
TITLE PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND  
OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION  
JOURNAL Patent: WO 9832462-A 21 30-JUL-1998;

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RESULT 2
LOCUS      A90886      20 bp      DNA      linear      PAT 22-JAN-2000
DEFINITION Sequence 21 from Patent EP0855184.
ACCESSION A90886
VERSION A90886.1 GI:6739328
KEYWORDS  unidentified.
SOURCE      unidentified.
ORGANISM    unidentified.
REFERENCE 1 (bases 1 to 20)
AUTHORS Heeg, K.P. and Lipford, G.B.
TITLE Pharmaceutical composition comprising a polynucleotide and an
antigen especially for vaccination
JOURNAL Patent: EP 0855184-A 21 29-JUL-1998;
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)
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LOCUS      AX45553      20 bp      DNA      linear      PAT 06-JUL-2002
DEFINITION Sequence 30 from Patent WO0222809.
ACCESSION AX45553
VERSION AX45553.1 GI:21714621
KEYWORDS  synthetic construct.
SOURCE      synthetic construct.
ORGANISM    artificial sequences.
REFERENCE 1
AUTHORS Bauer, S., Lipford, G. and Wagner, H.
TITLE process for high throughput screening of cpg-based
immuno-agonist/antagonist
JOURNAL Patent: WO 0222809-A 30 21-MAR-2002;
Coley Pharmaceutical GmbH (DE)
FEATURES
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DEFINITION Sequence 8 from patent US 5534409.
ACCESSION I23456
VERSION I23456.1 GI:1603326
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE 1 (bases 1 to 21)
AUTHORS Groner, B., Gouilleux, F. and Wakao, H.
TITLE Cytokine regulated transcription factor
JOURNAL Patent: US 5534409-A 8 09-JUL-1996;
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LOCUS      ARI74601      100 bp      DNA      linear      PAT 17-DEC-2001
DEFINITION Sequence 59 from patent US 6307024.
ACCESSION ARI74601
VERSION ARI74601.1 GI:17914921
KEYWORDS
SOURCE      Unknown.
ORGANISM    Unclassified.
REFERENCE 1 (bases 1 to 100)
AUTHORS Novak, J.E., Presnell, S.R., Sprecher, C.A., Foster, D.C., Holly, R.D.,
Gross, J.A., Johnston, J.V., Nelson, A.J., Dillon, S.R. and
Hammond, A.K.
TITLE Cytokine zaiaphall Ligand
JOURNAL Patent: US 6307024-A 59 23-OCT-2001;
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Db 67 AGATTCTTAGGAATTC AATC 86
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LOCUS      ARI74602/c      100 bp      DNA      linear      PAT 17-DEC-2001
DEFINITION Sequence 60 from patent US 6307024.

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ACCESSION      AR174602
VERSION        AR174602.1  GI:17914922
KEYWORDS
SOURCE         Unknown.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 100)
AUTHORS       Novak,J.E., Presnell,S.R., Sprecher,C.A., Foster,D.C., Holly,R.D.,
              Gross,J.A., Johnston,J.V., Nelson,A.J., Dillon,S.R. and
              Hammond,A.K.
TITLE         Cytokine zalphall Ligand
JOURNAL       Patent: US 6307024-A 60 23-OCT-2001;
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LOCUS          AX047032
DEFINITION     Sequence 37 from Patent WO0068381.
ACCESSION      AX047032
VERSION        AX047032.1  GI:11876456
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       artificial sequences.
REFERENCE      1 (bases 1 to 100)
AUTHORS       Presnell,S.R., Foster,D.C., Hammond,A.K. and Lok,S.
TITLE         Cytokine receptor mouse zcytor10
JOURNAL       Patent: WO 0068381-A 37 16-NOV-2000;
              ZymoGenetics, Inc. (US)
FEATURES      Location/Qualifiers
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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Db 67 AGATTTCTAGGAATTC AATC 86

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AX047033/c
LOCUS          AX047033
DEFINITION     Sequence 38 from Patent WO0068381.
ACCESSION      AX047033
VERSION        AX047033.1  GI:11876457
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       artificial sequences.
REFERENCE      1 (bases 1 to 100)
AUTHORS       Presnell,S.R., Foster,D.C., Hammond,A.K. and Lok,S.
TITLE         Cytokine receptor mouse zcytor10
JOURNAL       Patent: WO 0068381-A 38 16-NOV-2000;

ACCESSION      AR174602
VERSION        AR174602.1  GI:17914922
KEYWORDS
SOURCE         Unknown.
ORGANISM       Unclassified.
REFERENCE      1 (bases 1 to 100)
AUTHORS       Novak,J.E., Presnell,S.R., Sprecher,C.A., Foster,D.C., Holly,R.D.,
              Gross,J.A., Johnston,J.V., Nelson,A.J., Dillon,S.R. and
              Hammond,A.K.
TITLE         Cytokine zalphall Ligand
JOURNAL       Patent: US 6307024-A 60 23-OCT-2001;
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LOCUS          AX280202
DEFINITION     Sequence 48 from Patent WO0177171.
ACCESSION      AX280202
VERSION        AX280202.1  GI:16607595
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       artificial sequences.
REFERENCE      1
AUTHORS       Sprecher,C.A., Novak,J.E., West,J.W., Presnell,S.R., Holly,R.D. and
              Nelson,A.J.
TITLE         Soluble zalphall cytokine receptors
JOURNAL       Patent: WO 0177171-A 48 18-OCT-2001;
              ZymoGenetics, Inc. (US)
FEATURES      Location/Qualifiers
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Db 67 AGATTTCTAGGAATTC AATC 86

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DEFINITION     Sequence 49 from Patent WO0177171.
ACCESSION      AX280203
VERSION        AX280203.1  GI:16607596
KEYWORDS
SOURCE         synthetic construct.
ORGANISM       artificial sequences.
REFERENCE      1
AUTHORS       Sprecher,C.A., Novak,J.E., West,J.W., Presnell,S.R., Holly,R.D. and
              Nelson,A.J.
TITLE         Soluble zalphall cytokine receptors
JOURNAL       Patent: WO 0177171-A 49 18-OCT-2001;
              ZymoGenetics, Inc. (US)
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AX365191  
LOCUS AX365191 100 bp DNA linear PAT 15-FEB-2002  
DEFINITION Sequence 43 from Patent W0200721.  
ACCESSION AX365191  
VERSION AX365191.1 GI:18696947  
KEYWORDS synthetic construct.  
SOURCE synthetic construct.  
ORGANISM artificial sequences.  
REFERENCE 1 Sprecher, C.A., Presnell, S.R., Gao, Z., Whitmore, T.E., Kuijper, J.L. and Maurer, M.F.  
AUTHORS  
TITLE Cytokine receptor zcytor17  
JOURNAL Patent: WO 0200721-A 43 03-JAN-2002;  
ZymoGenetics, Inc. (US)  
FEATURES Location/Qualifiers  
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Matches 20; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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LOCUS AX365192 100 bp DNA linear PAT 15-FEB-2002  
DEFINITION Sequence 44 from Patent W0200721.  
ACCESSION AX365192  
VERSION AX365192.1 GI:18696948  
KEYWORDS synthetic construct.  
SOURCE synthetic construct.  
ORGANISM artificial sequences.  
REFERENCE 1 Sprecher, C.A., Presnell, S.R., Gao, Z., Whitmore, T.E., Kuijper, J.L. and Maurer, M.F.  
AUTHORS  
TITLE Cytokine receptor zcytor17  
JOURNAL Patent: WO 0200721-A 44 03-JAN-2002;  
ZymoGenetics, Inc. (US)  
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RESULT 13  
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DEFINITION Sequence 30 from patent US 6153397.  
ACCESSION AR119638  
VERSION AR119638.1 GI:14102337  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 31)  
AUTHORS Wisniewski, N., Silver, G.M., Lo, K. Callies. and Brandt, K.S.  
TITLE Flea epoxide hydrolase proteins and uses thereof  
JOURNAL Patent: US 6153397-A 30 28-NOV-2000;  
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LOCUS AR169119 31 bp DNA linear PAT 17-DEC-2001  
DEFINITION Sequence 30 from patent US 6290958.  
ACCESSION AR169119  
VERSION AR169119.1 GI:17906878  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 31)  
AUTHORS Wisniewski, N., Silver, G.M., Lo, K. Callies. and Brandt, K.S.  
TITLE Anti-flea epoxide hydrolase antibodies and uses thereof  
JOURNAL Patent: US 6290958-A 30 18-SEP-2001;  
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LOCUS AR138229 58 bp DNA linear PAT 16-JUN-2001  
DEFINITION Sequence 46 from patent US 6197930.  
ACCESSION AR138229  
VERSION AR138229.1 GI:14479738  
KEYWORDS  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 58)



AUTHORS Sheppard, P.O. and Humes, J.M.  
 TITLE Adipocyte-specific protein homologs  
 JOURNAL Patent: US 6197930-A 46 06-MAR-2001;  
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Copyright (c) 1993 - 2002 Compugen Ltd.

OM nucleic - nucleic search, using sw model  
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Title: US-09-355-254F-19  
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Searched: 2054640 seqs, 14551402878 residues

Total number of hits satisfying chosen parameters: 995600

Minimum DB seq length: 0  
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Post-processing: Minimum Match 0%  
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Listing first 45 summaries

Database : GenEmbl :

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- 39: em\_htgo\_hum:\*
- 40: em\_htgo\_mus:\*
- 41: em\_htgo\_other:\*

score greater than or equal to the score of the result being printed,  
and is derived by analysis of the total score distribution.

# SUMMARIES

Result No.	Score	Match	Length	DB	ID	Description
1	20	100.0	20	6	A89798	A89798 Sequence 20
2	20	100.0	20	6	A90885	A90885 Sequence 20
3	20	100.0	20	6	AX455567	AX455567 Sequence
4	20	100.0	21	6	I39725	I39725 Sequence 12
5	20	100.0	21	6	I55842	I55842 Sequence 12
6	18	90.0	86	6	AR184059	AR184059 Sequence
7	18	90.0	86	6	AR203344	AR203344 Sequence
8	18	90.0	86	6	AR206966	AR206966 Sequence
9	16.4	82.0	22	6	AR062163	AR062163 Sequence
10	15.2	76.0	24	6	AR061864	AR061864 Sequence
11	15	75.0	19	6	AR043736	AR043736 Sequence
12	15	75.0	19	6	AR043737	AR043737 Sequence
13	15	75.0	19	6	I81940	I81940 Sequence 38
14	15	75.0	19	6	I81941	I81941 Sequence 39
15	13.8	69.0	27	3	AF251171354	AF251171354
16	13.8	69.0	27	3	AF251171854	AF251171854
17	13.8	69.0	27	3	AF251174354	AF251174354
18	13.8	69.0	65	6	AX485415	AX485415 Sequence
19	13.6	68.0	41	6	AX088050	AX088050 Sequence
20	13.6	68.0	51	6	AX115257	AX115257 Sequence
21	13.6	68.0	100	1	ECOFIMPHC	M11776 E.coli inve
22	13.4	67.0	20	6	AR208137	AR208137 Sequence
23	13.4	67.0	60	6	AR009405	AR009405 Sequence
24	13.4	67.0	90	3	PFASSRR	D17580 Plasmodium
25	13.4	67.0	90	6	AR022104	AR022104 Sequence
26	13.4	67.0	90	6	AR022120	AR022120 Sequence
27	13.4	67.0	90	6	E08156	E08156 Primer or p
28	13.2	66.0	28	6	I39716	I39716 Sequence 3
29	13.2	66.0	28	6	I55833	I55833 Sequence 3
30	13.2	66.0	35	6	A92293	A92293 Sequence 12
31	13.2	66.0	35	6	A92344	A92344 Sequence 12
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33	13.2	66.0	38	6	I15748	I15748 Sequence 14
34	13.2	66.0	41	6	AX088048	AX088048 Sequence
35	13.2	66.0	48	6	AX429820	AX429820 Sequence
36	13.2	66.0	83	12	AY096764	AY096764 Populus t
37	13.2	66.0	90	10	RAT11BHVSD	M77835 R.norvegicu
38	13	65.0	13	6	AX026535	AX026535 Sequence
39	12.8	64.0	20	6	AX296543	AX296543 Sequence
40	12.8	64.0	24	6	AX291910	AX291910 Sequence
41	12.8	64.0	27	3	AF251728S4	AF251731 Dicyrtoma
42	12.8	64.0	27	3	AF251738S4	AF251741 Bilobella
43	12.8	64.0	48	9	HS4308535	AJ308535 Homo sapi
44	12.8	64.0	49	6	A43525	A43525 Sequence 5
45	12.8	64.0	49	6	AR052464	AR052464 Sequence

## ALIGNMENTS

RESULT 1	A89798	20 bp	DNA	linear	PAT 22-JAN-2000
LOCUS	Sequence 20 from Patent WO9832462.				
DEFINITION	A89798				
ACCESSION	A89798				
VERSION	A89798.1	GI:6738312			
KEYWORDS	unidentified.				
SOURCE	unidentified.				
ORGANISM	unclassified.				
REFERENCE	1 (bases 1 to 20)				
AUTHORS	Lipford,G.B. and Heeg,K.				
TITLE	PHARMACEUTICAL COMPOSITIONS COMPRISING A POLYNUCLEOTIDE AND				
JOURNAL	OPTIONALLY AN ANTIGEN ESPECIALLY FOR VACCINATION				
	Patent: WO 9832462-A 20 30-JUL-1998;				

Pred. No. is the number of results predicted by chance to have a

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LIPFORD GRAYSON B (DE); HEEG KLAUS (DE)  
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A90885  
LOCUS 20 bp DNA linear PAT 22-JAN-2000  
DEFINITION Sequence 20 from Patent EP0855184.  
ACCESSION A90885  
VERSION A90885.1 GI:6739323  
KEYWORDS unidentified.  
SOURCE unidentified.  
ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Heeg,K.P. and Lipford,G.B.  
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination  
JOURNAL Patent: EP 0855184-A 20 29-JUL-1998;  
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)  
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ORGANISM unidentified.  
REFERENCE 1 (bases 1 to 20)  
AUTHORS Heeg,K.P. and Lipford,G.B.  
TITLE Pharmaceutical composition comprising a polynucleotide and an antigen especially for vaccination  
JOURNAL Patent: EP 0855184-A 20 29-JUL-1998;  
HEEG KLAUS PROF DR (DE); LIPFORD GRAYSON B DR (DE)  
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LOCUS 20 bp DNA linear PAT 06-JUL-2002  
DEFINITION Sequence 44 from Patent WO0222809.  
ACCESSION AX455567  
VERSION AX455567.1 GI:21714635  
KEYWORDS synthetic construct.  
SOURCE synthetic construct.  
ORGANISM artificial sequences.  
REFERENCE 1  
AUTHORS Bauer,S., Lipford,G. and Wagner,H.  
TITLE Process for high throughput screening of cpq-based immuno-agonist/antagonist  
JOURNAL Patent: WO 0222809-A 44 21-MAR-2002;  
Coley Pharmaceutical GmbH (DE)  
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DEFINITION Sequence 12 from patent US 5648217.  
ACCESSION I55842  
VERSION I55842.1 GI:2476636  
KEYWORDS Unknown.  
SOURCE Unknown.  
ORGANISM Unclassified.  
REFERENCE 1 (bases 1 to 21)  
AUTHORS Levy,D.E.  
TITLE DNA sequence which binds transcriptional regulatory proteins activated in response to various cytokines and uses thereof  
JOURNAL Patent: US 5648217-A 12 15-JUL-1997;  
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DEFINITION Sequence 3 from patent US 6342581.

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ACCESSION AR184059
VERSION AR184059.1 GI:20228028
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 86)
AUTHORS Rosen,C.A., Ruben,S.M., Olsen,H.S. and Ebner,R.
TITLE Secreted protein HLHP03
JOURNAL Patent: US 6342581-A 3 29-JAN-2002;
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DEFINITION Sequence 6 from patent US 6365369.
ACCESSION AR203344
VERSION AR203344.1 GI:21499709
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 86)
AUTHORS Endress,G.A. and Rosen,C.A.
TITLE Prostate specific secreted protein
JOURNAL Patent: US 6365369-A 6 02-APR-2002;
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RESULT 8
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DEFINITION Sequence 18 from patent US 6372473.
ACCESSION AR206966
VERSION AR206966.1 GI:21505728
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 86)
AUTHORS Moore,P.A., Ruben,S.M. and Ebner,R.
TITLE Tissue plasminogen activator-like protease
JOURNAL Patent: US 6372473-A 18 16-APR-2002;
FEATURES
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DEFINITION Sequence 24 from patent US 5843697.
ACCESSION AR062163
VERSION AR062163.1 GI:5989854
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 22)
AUTHORS Pestka,S. and Kotenko,S.V.
TITLE Cells expressing IL-10 receptor and the CRFB4 gene product, an
JOURNAL Patent: US 5843697-A 24 01-DEC-1998;
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ACCESSION AR061864
VERSION AR061864.1 GI:5989555
KEYWORDS
SOURCE Unknown.
ORGANISM Unclassified.
REFERENCE 1 (bases 1 to 24)
AUTHORS Schumm,J.W., Micka,K.A. and Rabbach,D.R.
TITLE Multiplex amplification of short tandem repeat loci
JOURNAL Patent: US 5843660-A 56 01-DEC-1998;
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DEFINITION Sequence 106 from patent US 5814517.
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KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 19)
AUTHORS      Seidel,H.Martin., and Lamb,I.Peter.
TITLE        DNA spacer regulatory elements responsive to cytokines and methods
              for their use
JOURNAL      Patent: US 5814517-A 106 29-SEP-1998;
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VERSION       I81940.1  GI:3210237
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 19)
AUTHORS      Seidel,H.Martin., and Lamb,I.Peter.
TITLE        DNA spacer regulatory elements responsive to cytokines and methods
              for their use
JOURNAL      Patent: US 5814517-A 107 29-SEP-1998;
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Db 5 GATTTCCTCCCGAAATG 19

RESULT 12
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DEFINITION    Sequence 107 from patent US 5814517.
ACCESSION     AR043737
VERSION       AR043737.1  GI:5964745
KEYWORDS
SOURCE        Unknown.
ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 19)
AUTHORS      Seidel,H.Martin., and Lamb,I.Peter.
TITLE        DNA spacer regulatory elements responsive to cytokines and methods
              for their use
JOURNAL      Patent: US 5814517-A 107 29-SEP-1998;
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Db 5 GATTTCCTCCCGAAATG 5

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ACCESSION     I81940
VERSION       I81940.1  GI:3210237
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ORGANISM      Unclassified.
REFERENCE     1 (bases 1 to 19)
AUTHORS      Seidel,H.Martin., Lamb,I.Peter., and Chan,S.-S.Tian.
TITLE        Methods for detecting modulators of cytokine action
JOURNAL      Patent: US 5712094-A 38 27-JAN-1998;
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ACCESSION     I81941
VERSION       I81941.1  GI:3210238
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SOURCE        Unknown.
ORGANISM      Unknown.
REFERENCE     1 (bases 1 to 19)
AUTHORS      Seidel,H.Martin., Lamb,I.Peter., and Chan,S.-S.Tian.
TITLE        Methods for detecting modulators of cytokine action
JOURNAL      Patent: US 5712094-A 39 27-JAN-1998;
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DEFINITION    Acерentomon sp. CFNDS1 elongation factor 1-alpha gene, exon 4.
ACCESSION     AF251716
VERSION       AF251716.1  GI:10798923
KEYWORDS
SEGMENT       4 of 5
SOURCE        Acерentomon sp. CFNDS1.
ORGANISM      Acерentomon sp. CFNDS1.
              Eukaryota; Metazoa; Arthropoda; Hexapoda; Protura; Acерentomidae;
              Acерentomidae; Acерentomon.
REFERENCE     1 (bases 1 to 27)
AUTHORS      Carapell,A., Frati,F., Nardi,F., Dallai,R. and Simon,C.
TITLE        Molecular phylogeny of the Apterygotan insects based on nuclear and
              mitochondrial genes
JOURNAL      Pedobiologia (Jena) 44, 361-373 (2000)
REFERENCE     2 (bases 1 to 27)
AUTHORS      Carapell,A., Frati,F., Nardi,F., Dallai,R. and Simon,C.
TITLE        Direct Submission
JOURNAL      Submitted (30-MAR-2000) Evolutionary Biology, University of Siena,
              via P.A. Mattioli 4, Siena 53100, Italy
COMMENT       Region: Presence of introns of unknown length between s 162, 263,
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